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USAF SHALE OIL TO FUELS Volume II. Phases III and IV

UOP INC.
ALGONQUIN AND MT. PROSPECT ROADS
DES PLAINES, ILLINOIS 60016

JULY 1982

FINAL REPORT FOR PERIOD 1 OCTOBER 1980 - 31 DECEMBER 1981

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The novel hydrocracking scheme was successful in producing high quality jet fuel samples. A comparison of some key product properties with the specifications shows:

•	JP-4 JP-8		8	
•	Spec.	Product	Spec.	Product
Sp. Gr. 60/60°F	0.751-0.802	0.7813	$\overline{0.775} - 0.840$	0.7972
Smoke Point's	20	28.5	25	27.2
Aromatics, vol-%.	25	8.7	25	9.3
Combustion, Btu/lb	18,400	18,700	18,400	18,600

Based on yields and operating conditions demonstrated in pilot plant operations, process designs were prepared for the first- and second-stage hydrotreaters and the hydrocracker. These designs along with the information for the other process units were incorporated into a linear program model for a shale oil refinery. Using the basis provided by the USAF and assuming 100,000 barrels per stream day (BPSD) of raw shale oil valued at \$40/B, the following results were obtained:

	Max. JP-4	Max. JP-8
Jet Fuel Yield, BPSD	91,760	80,742
Total Liquid Product Yield	91,760	89,591
Capital Investment, MM \$ Product Cost, \$/B of Total	902	939
Liquid Product	56.82	58.80

FOREWORD

The final report describes the work completed by UOP Inc. on Phases III and IV of the Contract No. F33615-78-C-2079 entitled "United States Air Force Shale Oil to Fuels."

First Lieutenant Edward N. Coppola was the project engineer for the Air Force Wright Aeronautical Laboratories.

The work reported herein was performed during the period 1 October 1980 - 31 December 1981 under the direction of A. O. Braun and J. R. Wilcox. This report was submitted to the USAF in April, 1982.

The following individuals contributed to the execution of the program and the preparation of the report material:

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LIST OF SYMBOLS

Symbols Aromatics in FIA Analysis A Atomic Absorption Spectroscopy AAS Barrel В BFW Boiler Feed Water BPCD Barrels per Calendar Day Barrels per Stream Day BPSD British Thermal Units per Pound Btu/1b Benzene, Toluene, Xylene BTX °C Degrees Celsius Combined Feed Ratio С Combined Feed Ratio of Base Operation $C_{\mathbf{B}}$ Combined Feed Ratio, Vol. Feed Rate/(Feed Rate + Recycle CFR Rate) Centistoke cSt Discounted Cash Flow DCF Fouling Rate dR_F/dt Estimated Erected Cost EEC Engineering for Cost Estimating **EFCEST** End Point of Distillation EP U.S. Environmental Protection Agency EPA ۰F Degrees Fahrenheit FIA Fluorescent-Indicator Adsorption Gallon gal Gas Chromatograph GC Heat Transfer Coefficient h H₂ Recycle Ratio in SCFB H H₂ Recycle Ratio of Base Operation $H_{\mathbf{B}}$ High Pressure (Steam) HP K Degrees Kelvin kWh Kilowatt Hour Liquid Hourly Space Velocity, LHSV L LHSV of Base Operation L_{B} LHSV Liquid Hourly Space Velocity LP Low Pressure (Steam) Linear Programming LP MON Motor Octane Number Milliliter mL

Millions of U.S. Dollars Medium Pressure (Steam)

MM \$

MP

LIST OF SYMBOLS

Symbols	
N	Naphthenes in FIA Analysis
0	Olefins in FIA Analysis
P P P _B ppm psig	Paraffins in FIA Analysis Pressure Pressure of Base Operation Parts per Million Pressure, Pounds per Square Inch Gauge
Q	Heat Input
R _F RON RONC R+M/2 RVP	Fouling Factor Research Octane Number Research Octane Number Clear (no lead addition) Average of Research and Motor Octane Numbers Reid Vapor Pressure
SCFB SCFD SEM Sp. Gr. ST/D SUS	Standard Cubic Feet per Barrel Standard Cubic Feet per Day Scanning Electron Microscope Specific Gravity Short Tons per Day Saybolt Universal Seconds
T TB TEL TF FW	Temperature Temperature of Base Operation Tetra-ethyl Lead Fluid Temperature Wire Temeprature
vo1-%	Volume Percent

Weight Percent

wt-%

SUMMARY

The overall objective of this project was to demonstrate innovative technology to reduce the cost of converting shale oil to high yields of aviation turbine fuels. To carry out this program, UOP selected a processing scheme involving two stages of hydrotreating followed by hydrocracking. The Phase III and IV programs included first-stage hydrotreating catalyst stability testing, the production of jet and diesel fuel samples using the hydrocracking process, naphtha hydrotreating and catalytic reforming, arsenic management studies, an investigation of shale oil fouling, a shale oil/petroleum stability/compatibility study, and an economic analysis of the proposed upgrading scheme.

Based on the results of a six-month, first-stage hydrotreating operation, it was concluded that the catalyst showed excellent stability for arsenic and iron removal, and that high concentrations of contaminants could be contained. Fouling noted in the pilot plant preheater must be considered in a commercial design.

The novel hydrocracking scheme was successful in producing high quality jet fuel samples. A comparison of some key product properties with the specifications shows:

	JP-4		JP-8	
	Spec.	Product	Spec.	Product
Sp. Gr. 60/60°F	0.751-0.802	0.7813	0.775-0.840	0.7972
Smoke Point	20	28.5	25	27.2
Aromatics, vol-%	25	8.7	25	9.3
Combustion, Btu/1b	18,400	18,700	18,400	18,600

Catalytic reforming of the naphtha produced in the hydrocracking operation showed this material to be comparable to petroleum feedstocks and give C_5 plus liquid yields of 85 and 70 vol-% at 89 and 104 Research octane numbers (clear), respectively.

The aqueous solubility of arsenic on the used hydrotreating catalyst was found to exceed the acceptable level in the EPA Toxicity test. The arsenic-containing species were identified, thermal and chemical passivation and extraction techniques were investigated, and recommendations for further work were formulated.

Fouling studies showed that freshly produced samples would probably be much more reactive than the samples tested and that hydrotreating decreases the fouling tendency. A study of the compatibility and stability of blends of shale oils and petroleum showed no serious problems.

Based on yields and operating conditions demonstrated in pilot plant operations, process designs were prepared for the first- and second-stage hydrotreaters and the hydrocracker. These designs along with the information for the other process units were incorporated into a linear program

model for a shale oil refinery. Using the basis provided by the USAF and assuming 100,000 barrels per stream day (BPSD) of raw shale oil valued at \$40/B, the following results were obtained:

	Max. JP-4	Max. JP-8
Jet Fuel Yield, BPSD	91,760	80,742
Total Liquid Product Yield	91,760	89,591
Capital Investment, MM \$ Product Cost, \$/B of Total	902	939
Liquid Product	56.82	58.80

SECTION I

INTRODUCTION

This study was conducted to develop yield and economic data for the conversion of shale oil to aviation turbine fuels using innovative processing techniques. A processing scheme maximizing the yield of jet fuels at minimum cost was developed during Phase I of this program. This scheme consisted of the primary shale oil upgrading and conversion units with the necessary auxiliary facilities. The primary units include two stages of upgrading utilizing proprietary RCD Unibon® technology combined with UOP Hydrotreating technology. The upgraded shale oil is then converted to high yields of jet fuel using a novel, "parallel-flow" hydrocracking operation included in the UOP proprietary HC Unibon® technology.

The UOP approach to the problem of shale oil conversion to high quality fuels involves three distinct hydroprocessing steps. Shale oil has unique characteristics relative to conventional crude petroleum oil. Unusually high arsenic and iron levels, high pour point and viscosity, a high unsaturates concentration, and a significant solids (ash) content make conventional front-end refining techniques unusable without proper pretreatment.

The initial step in the UOP scheme is a feed preparation procedure which is essentially a desalting operation. The pilot plant utilized for this operation is completely analogous to a two-stage desalting unit commonly used in the refining industry.

After desalting, the first hydroprocessing step involves the use of UOP's RCD Unibon technology to effect metals removal, sulfur reduction, a degree of diolefin and olefin saturation, and the final solids clean-up necessary to render the resulting effluent suitable for subsequent processing.

Another characteristic of shale oil is its high nitrogen content relative to conventional crude oil. Reducing this contaminant to acceptable levels requires the use of a second-stage, high severity hydrotreatment of the first-stage hydrotreater effluent.

Once the metals, other contaminants and nitrogen contents have been reduced to acceptable levels and the unsaturates hydrogenated, shale oil is ready for the primary conversion step -- hydrocracking to jet and other fuels. The hydrocracking process that UOP has selected is a parallel-flow hydrocracker developed for conventional petroleum use.

Although jet fuel is the primary product in the overall refinery developed by UOP, diesel fuel and naphtha can also be produced. The naphtha was split into light and heavy fractions, with the heavy fraction converted to a gasoline component by reforming via a UOP Platforming® process unit. The reformate was mixed with the light naphtha fraction and sufficient butanes to produce the finished gasoline.

Phase I of the program provided for the development of innovative processing techniques and associated information for the conversion of shale oil to high yields of jet fuel. Phase II was primarily a test program utilizing small scale pilot plants to demonstrate the processing techniques developed in Phase I. The following summarizes the specific objectives for Phase II:

- Characterization of feedstock
- Determination of the shale oil fouling characteristics
- Preparation of sufficient feedstock to explore parallel-flow hydrocracking operating parameters
- Exploration of first- and second-stage hydrotreating operating parameters and catalyst systems
- Production of small scale jet fuel product samples

- Verification and/or adjustment of the operating requirement and product quality estimates generated during Phase I
- Establishment of processing parameter information for the Phase IV Economic Evaluation Study

Two feedstocks were used in this phase of the project. The first was shale oil derived from Occidental Petroleum Company's Modified In-Situ retorting process, and the second was shale oil obtained from the above-ground Paraho direct-heated retort. This work was summarized in Report Number AFWAL-TR-81-2116 entitled "United States Air Force Shale Oil to Fuels -- Phase II Interim Report" issued in November, 1981.

The objectives of the Phase III program included:

- Demonstration of first-stage hydrotreater catalyst stability
- Production of 5-gallon samples of jet and diesel fuels using the parallel-flow hydrocracking scheme
- Demonstration of hydrocracker catalyst stability
- Production of a 5-gallon sample of Special Grade gasoline using the Platforming process
- Investigation of crude shale oil arsenic solubilization and passivation or extraction of arsenic from used hydrotreating catalysts
- Determination of the relative fouling characteristics of raw and hydroprocessed shale oil fractions
- Examination of the stability and compatibility of shale oil and petroleum blends

The objectives of the Phase IV program included:

- Production of cost estimate based on process designs for the first-stage hydrotreater, second-stage hydrotreater and hydrocracker.
- Development of a shale oil refinery model
- Analysis of the economics of jet fuel production from raw shale oil

The results of the Phase III and Phase IV programs are contained in this final report.

The various operating conditions used during the pilot plant studies are consistent with commercial operating experience based on petroleum derived oils. All of the data presented in this report have operating conditions reported relative to a base operating condition. In all cases, the base conditions selected are those required to process a typical Middle East petroleum of the same boiling range to the identical product requirements. Positive differences in operating conditions indicate more severe requirements; negative differences indicate less severe processing requirements.

SECTION II

FIRST-STAGE HYDROTREATING

A six-month, first-stage hydrotreating run was made to provide additional processing and catalyst stability data required to establish the equipment design for a commercial plant. The specific data needed for this design work were the effect of metals deposition, particularly arsenic and iron, on catalyst activity and stability, when accomplishing the objective of reducing the arsenic content to less than 1 ppm.

Pilot Plant Description

A schematic diagram of the pilot plant used for this run is shown in Figure 1. Fresh feed and hydrogen are combined and flow concurrently downflow over the catalyst. The reactor effluent passes to a series of separators where the gas (mainly hydrogen) is separated, water scrubbed and recycled back to the reactor together with makeup hydrogen. The liquid is sent to a stripper to remove hydrogen sulfide and ammonia and is then collected under nitrogen in a glass receiver.

Based on the previously completed experimental work, the catalyst designated DRA was selected for this run. This proprietary, commercially-proven catalyst was developed to hydrotreat feeds containing large amounts of metal contaminants.

Feedstock

The feedstock was an Occidental shale oil produced from a modified in-situ retort. The "as-received" shale oil was dewatered and desalted in a two-stage electric desalter. Inspections of the treated shale oil are shown in Table 1. This 22.9 API gravity material contained 1.51 wt-% nitrogen, 27.5 ppm arsenic, and 45 ppm iron.

This feedstock was subjected to inspection throughout the six-month operation. As indicated on Table 2, the API gravity, distillation, and Bromine number show the expected variations in the test results. The values for arsenic show a broad range with many points below the 27.5 ppm level given in Table 1. The 60 ppm iron values in Table 2 are fairly consistent and higher than the 45 ppm level of Table 1.

Operating Conditions

Process conditions used in this study are comparable to those commercially employed for hydrotreating coke-oven light oil. The nomenclature adopted to relate the actual operating conditions to these base conditions is as follows:

		Written as
Temperature (Catalyst Average)	T-T (base), °F	$T-T_B$
Pressure	P-P (base), psi	$P-P_B$
Space Velocity	LHSV/LHSV base	L/L_{B}
H ₂ Recycle	SCFB/SCFB (base)	н/н _в

The start-of-run operating conditions were as follows:

Temperature	$T-T_B$, °F	50
Pressure	P-P _B , psi	0
Space Velocity	L/L _B	0.33
H ₂ Recycle	H/H _R	1.33

Stability Study Discussion

At the operating conditions given above, the catalyst maintained its activity for arsenic removal for the entire 170-day run. As shown in Figure 2, the product arsenic content was maintained at less than 0.2 ppm with no increase in catalyst temperature.

While achieving the desired arsenic removal level, the feed iron content was reduced to about 0.5 ppm for the entire run with no obvious

tendency to increase. The sulfur removal activity was also relatively constant giving a product value of about 0.05 wt-% for 92% removal.

A detailed, composite product analysis from late in the run is shown in Table 1. Product analyses completed at points throughout the run are given in Table 3. These data show that oxygen removal was relatively constant at about 50% whereas the nitrogen removal activity decreased from about 47% at start-of-run to about 22% at end of run. The conversion of the Conradson carbon components was relatively constant at 67%.

The overall yields for this operation are given in Table 4 and indicate a C_6 plus product in the range of 99 wt-% of fresh feed. About 580 standard cubic feet per barrel (SCFB) of hydrogen were required to accomplish this task.

At the end of the run, the catalyst was unloaded in five sections. It was free flowing and showed no signs of fouling. The used catalyst analysis given in Table 5 shows the pattern of higher to lower concentration of metal deposition from catalyst inlet to outlet. About 8.5 wt-% of arsenic was found on the chloroform-extracted catalyst removed from the top of the bed. The arsenic profile indicates that even after this sixmonth operation, the RCD catalyst had not reached its metals-limited capacity.

Fouling did occur three times during the course of the run. Each time, a deposit was formed at the top of the quartz chip preheater zone which caused a high pressure drop across the reactor, resulting in a "forced" shutdown. The reactor was opened and the fouled material removed and replaced. No loss in catalyst activity was observed as a result of these shutdowns.

The composition of the fouled material removed from the preheater in these three instances was quite similar. The data in Table 6 show that major constituents found on the quartz chips were carbon, hydrogen, arsenic, iron and sulfur, the remaining being trace metals. The presence

of these components is indicative of more than one fouling mechanism. These data also indicate that a successful commercial unit must have special features to accommodate these foulants.

The solids present in the desalted Occidental shale oil were subjected to additional analysis. A scanning electron microscope (SEM) was used to characterize the residue trapped on 5 micron filters and the toluene insoluble solids. As shown in the photographs of Figure 3 and 4, the material ranged in size from 5 to 200 microns, with the majority of the particles in the 10-20 micron range. Residue morphologies varied. The two most prevalent forms being agglomerated particles (often comprised of dense, flat platelets) and smooth dense individual particles exhibiting rounded edges.

The elemental analysis of the ash derived from these solids is given in Table 7. Using the emission spectrographic technique, analyses were completed on both the dry ash and the wet ash (sample is digested in sulfuric acid before ashing). Based on the large difference in the iron content, which is reported as ppm of metal in the oil, some of the iron-containing compounds may have been volatilized in the dry ash procedure. There are some inconsistencies in these data which remain unexplained and deserve further analysis. It can be concluded, however, that there is a wide size range of solids and that iron, nickel, sodium, calcium, magnesium and zinc are important components.

Conclusions

Based on the results of the six-month, first-stage hydrotreating operation, it is concluded that:

- 1. The UOP catalyst maintained its activity and stability for arsenic and iron removal over the entire 170-day run.
- 2. A total of 30 wt-% metals was accumulated on the catalyst during the course of the run.

- 3. Product arsenic was consistently below 1 ppm and iron at 1 ppm or less.
- 4. The fouling which occurred in the pilot plant preheater zone must be considered in the design of a commercial unit.
- 5. Additional experimental work to characterize the solids and investigate the fouling mechanisms is recommended.

TABLE 1. FIRST-STAGE HYDROTREATING OCCIDENTAL SHALE OIL

Feed and Product Comparison

	Feed	Product
API Gravity at 60°F	22.9	26.7
Specific Gravity at 60°F	0.9165	0.8944
Distillation (D-1160), °F		
IBP	376	370
5%	467	472
10%	570	508
30%	670	600
50%	712	698
70%	820	799
90%	953	940
95%	-	984
EP	-	-
% Over	87	92
Pour Point, °F	+65	+75
Viscosities		
Kinematic at 122°F, cSt	21.94	11.54
Kinematic at 210°F, cSt	5.268	3.473
Carbon, wt-%	84.85	84.85
Hydrogen, wt-%	12.27	12.63
Total Nitrogen, wt-%	1.51	1.17
Sulfur, wt-% (ppm)	0.64	(479)
Chloride, wt-ppm	< 1.0	-
BS and W, vol-%	0.2	-
Conradson Carbon, wt-%	1.36	0.45
Ash, wt-%	0.014	0.001
Heptane Insolubles, wt-%	0.34	0.01
Pentane Insolubles, wt-%	1.65	0.07
Metals by Emission (AAS), ppm		
Fe	45	(0.2)
N1	6.7	(2.2)
V	0.42	(< 0.1)
Pb	< 0.1	
Cu	< 0.1	
Na	11	
Мо	1.6	
Arsenic, wt-ppm	27.5	< 0.1
Bromine No.	23.6	7.5
Oxygen, wt-%	0.65	0.34
Water, wt-%	0.05	

TABLE 2. PERIODIC FEEDSTOCK ANALYSIS DURING STABILITY DEMONSTRATION

Period	1	95	138	198	246	270	318
Hours on Stream	Startup	1128	1654	2376	2952	3240	3816
API at 60°F	22.7	23.3	23.2	22.5	22.6	23.0	23.2
Sp. Gr. 60/60°F	0.9176	0.9141	0.9147	0.9188	0.9182	0.9159	0.9147
Distillation							
(D-1160), °F							
IBP	393	378	374	370	365	400	394
5%	478	459	468	458	480	482	488
10%	509	503	502	500	508	510	522
20%	570	569	560	560	559	561	572
30%	620	629	619	610	608	619	622
40%	675	688	675	659	660	680	672
50%	722	739	731	712	714	735	722
60%	770	789	781	771	769	787	78 0
70%	823	835	830	830	821	834	830
80%	882	880	881	883	871	882	890
85%				910		909	959
87%					930		
88%	941						
90%		979					
% Over	88.0	90.0	85.0	85.0	87.0	85.0	90.0
% Bottoms	12.0	10.0	15.0	15.0	13.0	15.0	10.0
Bromine Number	25.6	27.0	20.0	19.8		19.8	21.1
Arsenic, ppm	14.8	15.9	12.0	16.5	13.3	24	29
Iron, ppm	59.2	60	54	58	60	62	60
Ash, ppm		0.017		0.011			

TABLE 4. SINGLE-STAGE HYDROTREATER OCCIDENTAL SHALE OIL

Product Yields

Yields, Wt-% of Feed

н ₂ о	0.50
NH ₃	0.50
H ₂ S	0.65
c_1	0.03
c_2	0.06
c_3	0.09
C ₄	0.12
c ₅	0.15
c ₆ +	98.86
Total	100.96

Chemical Hydrogen Consumption = 581 SCFB

TABLE 5. FIRST-STAGE STABILITY DEMONSTRATION RUN

Used Catalyst Analyses

		Тор		Bottom	
	Top	Middle	<u>Middle</u>	<u>Miadle</u>	Bottom
Arsenic, wt-%	8.42	5.58	3.35	1.58	1.31
Iron, wt-%	6.00	3.74	1.91	0.99	0 .9 0
Carbon, wt-%	9.11	10.20	10.07	10.16	10.32
Sulfur, wt-%	6.20	6.38	5.66	5.77	5.51
Nitrogen, wt-%	0.39	0.43	0.43	0.85	0.22

TABLE 6. FIRST-STAGE STABILITY DEMONSTRATION RUN

Preheater and Plug Material

Time on Stream, Da	ıys		55		92	1	28
Location	•	Тор	Bottom	Top	Тор	Тор	Top
Dry Ash, wt-%		85.4	98.3	85.7	•	87.1	
Carbon, wt-%		10.4	0 .9 0	10.78	0.43	10.70	1.70
Hydrogen, wt-%		-	~	1.56		1.48	
Arsenic, AAS, wt-%	/	-		_	0.11	0.56	0.43
Sulfur, wt-%		-	-	-	-	1.52	-
Method:							
Emission Spec.	Quartz*						
•	Chips	Dry Ash	•		Wet Ash	Dry Ash	Wet Ash
Metal, wt-%	•	•				•	
Fe	0.044	3.4	1.2	1.2	0.96	2.4	2.6
Ni	0.004	0.064	0.052	<0.03	<0.03	0.12	0.13
V	-	< 0.003	<0.003	-	<0.03	_	< 0.03
Ca	<0.03	0.068	0.032	-	<0.2	<0.3	<0.2
Mg	0.006	0.010	0.005	0.015	0.012	< 0.01	0.015
Mn	0.002	0.012	0.009	0.015	0.020	0.015	0.014
Cr	0.002	0.039	0.007	0.038	0.098	0.025	0.03
Sn	_	0.010	<0.003	<0.03	< 0.03	<0.03	<0.03
Cu	<0.001	0.059	0.013	0.013	< 0.01	0.017	0.017
Zn	-	0.015	0.25	< 0.3	< 0.4	< 0 • 4	<0.4
Ti	0.16	0.18	0.18	0.32	0.58	0.30	0.35
Pb	-	0.003	< 0.001	<0.03	< 0.04	< 0.04	< 0.04
Na	< 0.05	0.45	0.025	< 1.0	< 1	< 0.5	< 1
Мо	_	0.13	0.059	0.043	< 0.04	0.11	0.16
Со	0.021	0.006	0.070	0.078	0.086	0.15	0.15
Si	Major						
Al	0.46	0.49	0.58	0.37	0.60	0.38	0.45
Zr	0.006	_	-				
В	0.03	-	-				
Ba	0.006	0.007	0.005				

^{*} This analysis is included to provide a point of reference.

TABLE 7. SOLIDS ANALYSIS

Desalted Occidental Shale Oll

Total		Filter Cake on	
Sample	วิบุm Filter	10µ m Filter	25 µm Filter
165			
stable			
rt−\$ 0.003			
Ash			
	**	**	**
	9.8	23.0	8.3
			0.04
			0.02
			0.01
			0.01
			0.07
			0.14
			0.03
			0.31
	0.00	0,01	0.51
	0.27	0.40	0.13
			0.09
			0.33
			1.9
			0.01
	0.02	0.05	0.01
2.0			
4.8			
	Sample 165 0.015 0.015 0.01	Sample Sum Filter	Sample Sum Filter 10µ m Filter

Мо

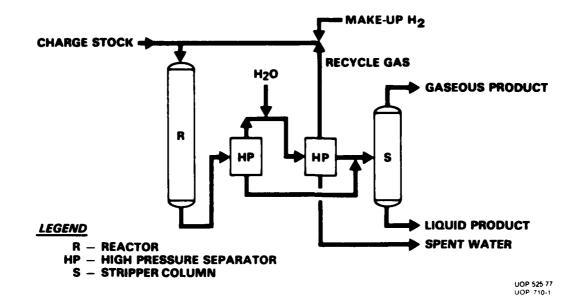


FIGURE 1

FIRST STAGE HYDROTREATING PILOT PLANT

SCHEMATIC FLOW DIAGRAM

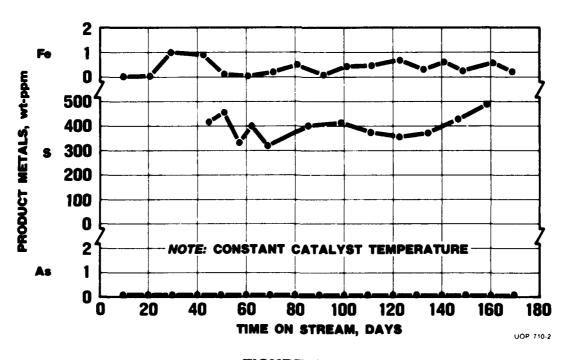
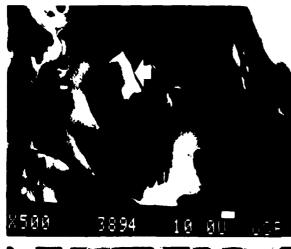
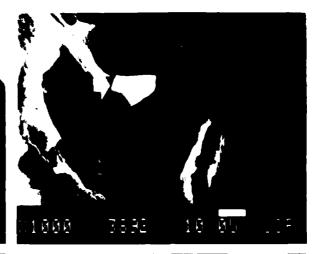


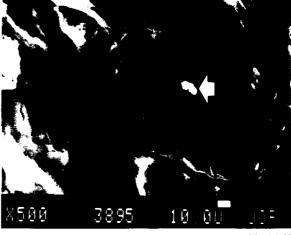
FIGURE 2

FIRST STAGE HYDROTREATING CATALYST STABILITY DEMONSTRATION









UOP 734-23

1...

FIGURE 3

SCANNING ELECTRON MICROSCOPE ANALYSIS OF OCCIDENTAL SHALE OIL RESIDUE

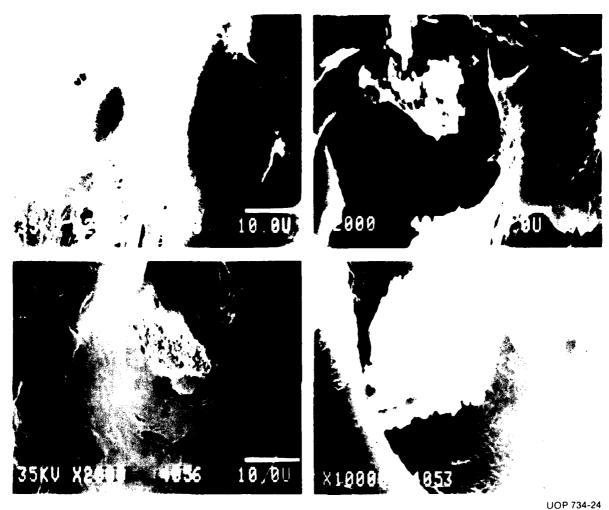


FIGURE 4

SECTION III

HYDROCRACKING

As part of the Phase III pilot plant program, three hydrocracking studies were completed: preparation of 5-gallon samples of jet and diesel fuels, determination of catalyst stability in the parallel-flow scheme, and investigation of sulfur addition to the hydrocracker feedstock. The data developed from these studies were also used to confirm the refinery design and provide other information needed in the economic evaluation program of Phase IV.

Pilot Plant Description

The pilot plant used in the hydrocracking operation embodies the essential features of a commercial unit. Fresh feed, recycle liquid, recycle gas and makeup hydrogen are charged to the reactor section. Trickle-bed reactors containing 25-200 mL of catalyst were employed. Gas is recycled from the high pressure separator back to the reactor section. The reactor liquid effluent is charged to a series of separators including a fractionator and debutanizer to produce liquid product and a recycle liquid stream. In the pilot plant operations, reactor temperatures were adjusted to achieve 100% conversion of the feed to products. During these operations, no fractionator bottoms product was withdrawn as a net product.

The DCC catalyst used in this study is a proprietary material which has seen wide commercial operation. This is the same catalyst which was used successfully in the Phase II hydrocracking programs.

Feedstock

The hydrocracker feedstock is a second-stage hydrotreated Occidental shale oil, whose properties are shown in Table 8. This stock was produced

by first hydrotreating the raw shale oil at relatively mild conditions to remove the contaminants and stabilize the material, and then hydrotreating the first-stage product at relatively severe conditions to reduce the high levels of impurities, such as nitrogen and oxygen, that were still present. The measured nitrogen content of the material was 780 ppm. The hydrocracker feed sulfur content was 139 ppm, the oxygen content 545 ppm, and the Bromine number 1.1.

Operating Conditions

The operating conditions used in the production of the fuel samples were in the range of those commercially practiced, and for this report, are presented relative to the conditions appropriate for converting a typical Arabian vacuum gas oil. The nomenclature adopted is as follows:

		Written as
Temperature (Catalyst Average)	T-T (base), °F	$T-T_B$
Pressure	P-P (base), psi	P-PB
Space Velocity	LHSV/LHSV (base)	L/L _B
Combined Feed Ratio	CFR/CFR (base)	c/c _B
H ₂ Recycle	SCFB/SCFB base	н/н _в

The conditions in the two operations, Runs 1 and 2, were varied depending on the type of fuel being produced. Run 1 was designed to produce the jet fuel samples. The latter portion of the run was used to establish operating conditions and procedures for the diesel fuel production and the stability demonstration, in Run 2.

Run 2 proceeded initially in the JP-8 mode to obtain the base activity-conversion level. This was followed by the more severe diesel fuel production mode. The final processing was again in the JP-8 mode to obtain the final activity-conversion level and the catalyst deactivation rate.

Run		1				2	
Type of Fuel	JP-8	JP-4	Diesel	JP-8	Diesel	JP-8	JP-8
Hours on Stream	70-360	360-610	610-860	0-150	150-460	460-780	780-1293
Temperatures, (T-T _B), °F	- 54	-50	-44	-53	-49	-43	-45
Pressure, (P-P _B), psig	-150						
Space Velocity, (L/L_B)	1.4	1.4	2.0	1.4	2.0	1.4	1.4
CFR, (C/C _B)	1.0	1.0	0.8	1.0	0.8	1.0	1.0
H_2 Recycle, (H/H _B)	1.4						

Sample Production

Representative product yield distributions obtained when hydrocracking the hydrotreated Occidental shale oil to JP-4 and JP-8 fuels are given in Table 9. The JP-4 yield was better than 84 wt-% and required 1029 SCFB of hydrogen. The JP-4 volume yield amounted to about 93 vol-% of hydrocracker feed. The equivalent values for JP-8 are 75 wt-% yield (80 vol-%) with 921 SCFB hydrogen consumption. Both of these yield patterns show the excellent selectivity of the parallel-flow scheme for jet fuel production.

More detailed product distributions for the JP-4 and JP-8 cases are given in Tables 10 and 11, respectively. Both of these analyses are from single test periods and differ slightly from the average values given in Table 9. Included with the more detailed light ends yields are the iso-and normal-paraffin breakdown for the butanes and pentanes. About 60 wt-% of the butanes and 67 wt-% of the pentanes are in the more valuable non-normal configuration.

The UGP inspections of the JP-4 jet fuel sample are shown in Table 12 along with the USAF specifications. With the exception of the slightly high 50 and 90% points of the distillation and the conductivity, this

product meets all U.S. military specifications and should be environmentally acceptable in view of the low nitrogen and sulfur contents. With only 8.7 vol-% aromatics reported (25.0 vol-% is the maximum specification), the Smoke Point exceeds the minimum acceptable level of 20 by 8.5. The net combustion value is some 300 Btu/lb higher than required and is a reflection of the high hydrogen content obtained in the parallel-flow hydrocracking operation.

Inspections of the JP-8 jet fuel produced are shown in Table 13, along with the USAF specifications. These inspections meet all of the U.S. military specifications except for Freeze Point (-54°F vs. -58°F) and conductivity. These can be easily met by slightly lowering the end point and including an additive, respectively. This higher end point fuel, 552°F vs. 517°F for the JP-4, contains 9.7 vol-% aromatics and gave a Smoke Point of 27.2 (minimum specification is 25). The combustion value is 200 Btu/lb higher than required and again shows the high hydrogeration capability of the parallel-flow hydrogracking operation.

An analysis of the C_5 - C_6 fraction obtained during the JP-8 production run is given in Table 14. This material is highly isomerized and contains less than 1 wt-% benzene. The clear Research octane number of 70.6 increased to 87.7 with the addition of 3 cc TEL per gallon.

A detailed analysis of the C7-232°F naphtha yielded during the JP-8 production run is given in Table 15. This highly saturated material contains 96.8 vol-% paraffins plus naphthenes and has a clear Research octane number of 60.2. After treatment to remove the small quantity of sulfur remaining, this naphtha represents a good feedstock for further upgrading by catalytic reforming to produce both a high octane blending component and hydrogen for the shale oil hydrotreating operations. The catalytic reforming of this material is discussed later in this report.

The product distributions for hydrocracking the hydrotreated Occidental shale oil to DF-2 and DFM are shown in Table 16. The yields of 98.6 wt-% and 96.2 wt-% were achieved for DF-2 and DFM, respectively. Both fuels were produced with the same end point and, therefore, the yield

difference results from adjusting the amount of C_5 - C_6 material in the product to meet the specified flash points. As noted in the jet fuel production cases, the parallel-flow hydrocracking scheme showed excellent selectivity for converting the hydrotreated shale oil into these distillate fuels. The chemical hydrogen consumption was 800 SCFB for both the DF-2 and DFM cases.

A more detailed product distribution obtained in a single test period in the diesel fuel production mode is given in Table 17. The C_4 - yield in this operating mode is only about one-third that observed in the more severe jet fuel operations. The extent of butane and pentane isomerization is about the same: 60% iso in the total butane and 67% iso in the total pentane.

Inspections of the DF-2 and DFM samples, together with the U.S. military specifications, are shown in Tables 18 and 19, respectively. These inspections show that the diesel fuels met all specifications. Both samples had Cetane numbers greater than 55 and very low sulfur contents of about 1 ppm. Both diesel fuels show excellent stability and should be environmentally acceptable. In addition, their low acid numbers, copper strip corrosions, sediments, particulates, excellent stabilities and high cetane numbers should pose no problem for storage and use in engines.

Catalyst Stability Demonstration

A second objective in the Phase III hydrocracking program was to make an extended catalyst stability study to provide information regarding longer term effects of processing shale oil in the proposed UOP turbine fuels refinery and to assess the advantage of parallel-flow over single-stage hydrocracking in terms of catalyst stability.

As indicated previously, the stability run was made in the same hydrocracking operation used for the sample production. After checking the plant operation in the JP-8 mode for 150 hours, and then producing the diesel samples, the plant was switched back to the JP-8 mode, and the operation continued for a period of 1200 hours to measure the catalyst deactivation.

The deactivation was measured by the increase in reactor temperature per unit of operating time required to maintain constant conversion to a specified product end point. In this case, the end point was that required to make JP-8 jet fuel. Results of the run are shown in Figure 5. The rate of deactivation is slightly lower than was reported for a similar operation over a shorter period of time in the Phase II program and as shown in the Phase II report, much improved over the expected result for the single-stage operation.

During the catalyst stability demonstration (Run 2), and at the end of Run 1, plugging of the reactor preheater was observed. Elemental analyses of the material recovered from the inert, granular material (quartz chips) are given in Table 20. Relatively high concentrations of iron, nickel and chromium are found. Analyses of the carbonaceous material found at the same time periods on a filter in the fresh feed line, as given in Table 21, indicate the presence of high molecular weight, heteroatomic materials. The presence of these materials in the hydrotreated feedstock and their subsequent fouling of the reactor preheater are believed due to oxidation and degradation as a result of handling and storing this material during a 9-month period.

Sulfur Addition

In the Phase II hydrocracking program, it had been observed that the addition of sulfur to the relatively sulfur-free hydrocracker feedstock improved the activity of the catalyst. With the feedstock containing less than 3 ppm sulfur, it was susplicted that this was insufficient to maintain the catalyst in its activated (sulfided) state.

After the period in which the stability of the catalyst was demonstrated, 10,000 ppm of sulfur as ditertiary-butyl disulfide was added to the feed. As before, the catalyst temperature required to maintain conversion was reduced by about 10°F. The presence of plant operating problems in the fractionation section forced this study to be discontinued before a definitive response could be determined.

Conclusions

Based on the results of these hydrocracking operations, it was concluded that:

- 1. The parallel-flow hydrocracker can produce military turbine fuels and diesel fuel in excellent yields.
- These fuels not only meet, but exceed, military specifications in many areas.
- 3. The parallel-flow hydrocracker offers improved catalyst stability when compared with conventional single-stage hydrocracking.

TABLE 8. HYDROCRACKING FEEDSTOCK

Occidental Shale Oil

		Boiling P	t. GC, Wt-%
API Gravity at 60°F	32.0		
Sp. Gr., 60/60°F	0.8654	% Over	Temp., °F
Distillation (D-1160), vol-%		IBP	241
IBP, °F	390	5	361
5%	469	10	421
10%	495	15	463
30%	581	20	495
50%	663	25	519
70%	760	30	545
90%	898	35	571
95%	958	40	592
EP, °F	988	45	615
% Over	97. 0	50	640
		55	666
Carbon, wt-%	86.99	60	692
Hydrogen, wt-%	13.41	65	718
Nitrogen, ppm	78 0	7 0	747
Sulfur, ppm	139	75	776
Pentane Insolubles, wt-%	0.12	80	804
Heptane Insolubles, wt-%	0.05	85	832
Conradson Carbon, wt-%	0.0 9	9 0	872
Ash, wt-%	0.001	9 5	931
Bromine Number	1.1	FBP	1100
BS & W, wt-%	0.4		
Toluene Insolubles, wt-%	0.01		
Oxygen, ppm	545		
Molecular Weight	305		

TABLE 9. HYDROCRACKER PRODUCT DISTRIBUTIONS Hydrotreated Occidental Shale Oil Feed

Product Desired	JP-8	JP-4
Wt -%		
Fresh Feed	100.00	100.00
H_2 Consumption, wt-%	1.61	1.80
H ₂ Consumption, SCFB	921	1029
NH ₃	0.08	0.08
H ₂ S	0.01	0.01
c ₁ -c ₃	1.50	1.88
C ₄	3.73	4.80
c ₅ -c ₆	9.29	11.03
C ₇ -250°F	11.75	JP-4 12.18 71.82**
250-EP	JP-8 <u>75.25</u> *	71.82**
Total	101.61	101.80
<u>Vol-%</u>		
Fresh Feed	100.00	100.00
c ₄	5.78	7.43
c ₅ -c ₆	12.62	14.75
C ₇ -250 °F	13.76	JP-4 \begin{cases} 14.31 \\ 78.68***
250-EP	JP-8 <u>80.40</u> *	78.68**
Total	112.56	115.17

Product EP ∿550°F Product EP ∿520°F

TABLE 10. HYDROCRACKING HYDROTREATED OCCIDENTAL SHALE OIL

JP-4 Product Distribution

	Yield, Wt-%	Yield, Vol-%	Gas Yield, SCF/BBL
Hydrogen	0.23		
H ₂ S	0.01		
NH ₃	0.16		
Methane	0.24		16.4
Ethane	0.25		9.1
Propane	1.39		34.2
Isobutane	2.83	4.44	53.0
n-Butane	1.97	2.98	36.9
Isopentane	3.29	4.63	
n-Pentane	1.70	2.37	
Hexane	6.03	7.74	
C ₇ -250°F Fraction	12.17	14.29	
250-EP Fraction	71.74	78.59	
Total	102.01	115.03	149.7
Subtotals			
C ₁ -C ₄ Fraction	6.68		149.7
$C_5^1-C_6^2$ Fraction	11.02	14.73	
C_7^{-250} °F Fraction	12.17	14.29	
250-EP Fraction	71.74	78.59	
C ₅ + Fraction	94.93	107.61	
Percent Iso in			
Butane	58.94		
Pentane	65.85		

TABLE 11. HYDROCRACKING HYDROTREATED OCCIDENTAL SHALE OIL

JP-8 Product Distribution

	Yield, Wt-%	Yield, Vol-%	Gas Yield SCF/BBL
Hydrogen	0.22		
H ₂ S	0.01		
NHี ₃	0.16		
Methane	0.18		12.0
Ethane	0.18		6.5
Propane	1.09		26.9
Isobutane	2.42	3.81	45.4
n-Butane	1.52	2.29	28.4
Isopentane	3.02	4.26	
n-Pentane	1.42	1.97	
Hexane	4.98	6.60	
C ₇ -232°F Fraction	7.73	8.71	
232-EP Fraction	78.90	85.65	
Total	101.83	113.28	119.2
Subtotals			
C ₁ -C ₄ Fraction	5.39		119.2
C ₅ -C ₆ Fraction	9.42	12.83	
C ₇ -232°F Fraction	7.73	8.71	
232-EP Fraction	78.90	85.65	
C ₅ + Fraction	96.05	107.18	
Percent Iso in			
Butane	61.5		
Pentane	68.1		

TABLE 12. JP-4 JET FUEL

Production Sample

	USAF Specs.	
API Gravity at 60°F	45 - 57	49.6
Sp. Gr. 60/60°F		0.7813
Distillation (D-86), vol-%		0.7015
IBP, °F	report	202
5%	•	236
10%	report	255
20%	293	288
30%		321
40%		354
50%	374	386
60%		414
70%		437
80%		456
90%	473 max.	478
95%		495
EP, °F	518 max.	
	J 20 maze	517
Freeze Point, °F	-72 max.	Below -70°F
Smoke Point	20 min.	28.5
Vapor Pressure, 38°C, psi	2-3 max.	
Viscosity -20°C, cSt		0.8 3.207
Acid No., mg KOH/mg	0.015 max.	0.012
Copper Strip Corrosion	1B max.	
Coulometric Sulfur, ppm	0.4 wt-% max.	1A 737
Mercaptan Sulfur, wt-%	0.001 max.	0.0001
Coulometric N, ppm	30002	0.37
Carbon, wt-%		84.23
Hydrogen, wt-%	13.6 min.	14.39
FIA, vol-%	1300	14.37
P + N		01 2
0	5.0 max.	91.3
A	25.0 max.	8.7
Combustion, Btu/1b	18,400 min.	18,700
Existent Gum,	- · · · · · · · · · · · · · · · · · · ·	10,700
Unwashed/Washed, mg per 100 mL	7 max.	7.8/4.0
Naphthalenes, UV wt-%		0.28
Conductivity	50-300	
Water Separation D-1094		1 p̄ S/M
Interface Rating	1B max.	1B
Separation Rating	l max.	1
Demulsification		No Emulsion
		0il Layer Clear
		No Scum
JFTOT		ocum
m 0		

Temp., °C	Min.	ΔP mm Hg	TDR Spun	ASTM Code
260	150	1.5	1.5	0

TABLE 13. JP-8 JET FUEL

Production Sample

	USAF Specs.	
API Gravity at 60°F	37-51	46.0
Sp. Gr. 60/60 F	0.775-0.840	0.7972
Distillation (D-86), vol-%		
IBP, °F	report	288
5%	•	308
10%	401 max.	321
20%		352
30%		384
40%		412
50%		434
60%		455
70%		474
80%		492
90%		512
95%		525
EP, °F	572 max.	552
% Over		99. 0
Smoke Point	25 min.	27.2
Freeze Point, °F	-58 max.	- 54
Flash Point, °F	100 min.	100
Viscosity -20°C, cSt	8.0 max.	5.670
Acid No., mg KOH/gm	0.015 max.	0.01
Coulometric Sulfur, ppm	0.4 wt-% max.	499
Mercaptan Sulfur, wt-%	0.001 max.	0.00048
Carbon, wt-%		84.63
Hydrogen, wt-%	13.6 min.	14.14
FIA, vol-%		
P + N		90.7
0	5.0 max.	_
A .	25.0 max.	9.3
Combustion, Btu/1b	18,400 min.	18,600
Copper Strip Corrosion	1B max.	1A
Coulometric N, ppm		0.7
Existent Gum,	_	
Unwashed/Washed, mg per 100 mL	7 max.	4.1
Naphthalenes, UV wt-%	3 max.	0.35
Conductivity	50-300	4 p̄ S/M
Water Separation D-1094	4.00	
Interface Rating	1B max.	1
Separation Rating	2 max.	1

<u>JFTOT</u> <u>Temp., °C</u>	Min.	ΔP mm Hg	TDR Spun	ASTM Code
260	150	0	1.0	0

TABLE 14. C5-C6 FRACTION FROM JP-8 JET FUEL

API 60°F Sp. Gr., 60/60°F	77.2 0.6780	Composition Wt-%	-
Distillation (D-86), vol-%			
IBP, °F	124	1-C4	0.1
5%	134		0.2
10%	136	<u>n</u> -C ₄ 1-C ₅	5.8
20%	140	n-C5	6.3
30%	141	2, 2 DMB	0.4
40%	144	CP	0.6
50%	146	2,3 DMB	2.7
60%	148	2 MP	22.3
70%	150	3 MP	15.4
80%	152	n-C ₆	22.1
90%	156	MCP	17.2
95%	161	CH	2.1
EP, °F	220	Bz	0.9
% Over	99.0	C ₇ +	3.9
Coulometric S, ppm	1.0		
Nitrogen, ppm (Chem Lum)	0.83		
RON Clear	70.6		
Leaded (3 cc TEL/gal)	87.7		

TABLE 15. C7-232°F FRACTION FROM JP-8 JET FUEL

API at 60°F	64.0		
Sp. Gr., 60/60°F	0.7238		
D1 - (11 - (D 0())	g/	GC, Wt-%	
Distillation (D-86), vol		A A - P	
IBP, °F	194	Aromatics	2 5
5%	197	Toluene	3.5
10%	198	$\frac{m}{m}$ x 1	0.1
20%	199 201	Total	3.6
30%		Donaffina/Nanhahana	
40%	202	Paraffins/Naphthenes	2 02
50%	203	C ₆ Paraffins	2.83
60%	205	C ₆ Cyclics	2.74
70%	207	C7 Paraffins	44.38
80%	209	C7 Cyclics	28.34
90%	216	Cg Paraffins	10.43
95%	222	C ₈ Cyclics	8.39
* EP, °F	278	Co Paraffins	.19
% Over	99.0	C ₉ Cyclics	0.10
			97.40
Coulometric S, ppm	17.1		
Nitrogen, ppm (Chem Lum)	0.40		
Carbon, wt-%	83.76		
•			
Hydrogen, wt-%	16.00		
FIA, vol-%			
P + N	96.8		
0	-		
A	3.2		
RON Clear	60.2		
RON Leaded (3 cc TEL/gal	.) 82.2		

^{*} Dry Point = 232°F

TABLE 16. HYDROCRACKING HYDROTREATED OCCIDENTAL SHALE OIL TO DF-2 AND DFM DIESEL FUELS

Product Distribution, wt-%	Feed	DF-2	DFM
н ₂ s		0.01	0.01
NH ₃		0.10	0.10
н ₂ о		0.06	0.06
C ₄ minus		1.68	1.68
C ₅ and/or C ₆		1.00	2.36
Flash Point* to EP Diesel Fuel	44.0	98.55	96.19
Total		101.40	101.40
* Flash Point, °F		133	140
Hydrogen Consumption, SCF/B		80	00

TABLE 17. HYDROCRACKING HYDROTREATED OCCIDENTAL SHALE OIL

Diesel Product Distribution

	Yield, Wt-%	Yield, Vol-%	Gas Yield, SCF/BBL
Hydrogen	0.13		
H ₂ S	0.01		
NH3	0.10		
Methane	0.13		8.9
Ethane	0.11		4.0
Propane	0.38		9.3
Isobutane	0.68	1.07	12.7
n-Butane	0.48	0.72	8.9
Isopentane	0.82	1.16	
n-Pentane	0.40	0.56	
Hexane	1.30	1.72	
C ₇ -183°F Fraction	0.19	0.16	
183-EP Fraction	96.77	100.78	
Total	101.51	106.17	43.8
Subtotals			
C ₁ -C ₄ Fraction	1.77		43.8
C_5 - C_6 Fraction	2.53	3.34	
C ₇ -183°F Fraction	0.19	0.16	
183-EP Fraction	96.77	100.78	
C ₅ + Fraction	99.49	104.28	
Percent <u>Iso</u> in			
Butane	58.8		
Pentane	67.05		

TABLE 18. DIESEL FUEL DF-2

Production Sample

	USAF Specs.	
API Gravity at 60°F Sp. Gr. 60/60°F	32.9-41	38.8 0.8309
Distillation (D-86), vol-%		
IBP, °F		32 0
5%		362
10%		400
20%		452
30%		486
40%		513
50%	report	542
60%		568
70%		593
80%		618
90%	675 max.	646
95%		664
EP, °F	700 max.	678
% Over		98.5
Flash Point, °F	133 min.	134
Cloud Point, °F	report	32
Pour Point, °F		5
Aniline Point, °F		173.1
Viscosity 100°F, cSt	1.8-9.5	3.367
D-1500 Color	report	<2
Ac'd No., mg KOH/gm	0.10 max.	0.018
Copper Strip Corrosion, 100°C	l max.	1 A
Coulometric Sulfur, ppm	0.70 wt-% ma	1.2
Cetane Number	45 min.	55.1
Water and Sediment, D-2709, wt-%	0.01 max.	0.005
Demulsification, 25°C		No Emulsion
		011 Layer Clear
Particulate Cont. D-2276, mg/L	8 max.	8
Carbon Residue on 10% Botts,		
D-524, wt-%	0.2 max.	0.11
Ash, wt-%	0.02 max.	0.001
a. 1.11. p. 207/ /100 r		

1.5 max.

0.45

0.35

Total 0.8

Stability D-2274, mg/100 mL

Adherent Gum

Sediment

TABLE 19. DIESEL FUEL MARINE DFM

Production Sample

USAF Specs.

API Gravity at 60°F Sp. Gr. 60/60°F	report	38.7 0.8314	
Distillation (D-86), vol-%		2/0	
IBP, °F		348	
5%		376	
10%		420	
20%		465	
30%		498	
40%		525	
50%	report	550	
60%		575	
70%		598	
80%		624	
90%	675 max.	650	
95%		672	
EP, °F	725 max.	680	
% Over		99.0	
Flash Point, °F	140 min.	144	
Cloud Point, °F	30 max.	28	
Pour Point, °F	20 max.	5	
Aniline Point, °F	report	173.1	
Viscosity 100°F, cSt	1.7-4.3	3.444	
D-1500 Color	3 max.	<3	
Acid No., mg KOH/gm	0.30 max.	0.021	
Copper Strip Corrosion, 100°C	1 max.	1 A	
Coulometric Sulfur, ppm	1.0 wt-%	max. 1.9	
Cetane Number	45 min.	55.3	
Water and Sediment, D-2709, wt-%		<0.005	
Demulsification, 25°C, minutes	10 max.	No Emulsion	
, ,		Oil Layer Clea	ir
Particulate Cont. D-2276, mg/L		5	
Carbon Residue on 10% Botts,			
D-524, wt-%	0.2 max.	0.11	
Ash, wt-%	0.005 max.	<0.001	
Stability D-2274, mg/100 mL			
Adherent Gum	2.0 max.	0.3	
Sediment	aro many	0.3	Total 0.7
Coa Line is a		0.5	

TABLE 20. PREHEATER PLUG ANALYSES

Run 2

Time on Stream, hours		783	End of Run 1703
Plug Number		1	2
Solids, wt-%	Quartz Chips	96.2	92.5
Emission, wt-ppm	-		
Fe	440	260	870
Ni	40	30	70
V	-	30	60
Ca	300	280	290
Mg	60	10	80
Mn	20	20	30
Cr	20	-	-
Sn		-	-
Cu	10	50	140
Zn	-	-	40
Ti	1600	1500	1 9 00
Pb	_	-	-
Na	500	1000	1000
Мо	-	30	30
Со	210	~	-
Si	Ma jor	Ma jor	Ma jor
A1	4600	2300	5300
Ва	60	3 0	60
Sr	-	-	
Zr	60	240	310
В	300	800	110
Carbon, wt-%		53.67	6.90
Sulfur, wt-%		5.12	

TABLE 21. FRESH FEED FILTER CAKES

Run 2

Time on Stream, hours	783	End of Run 1703
Molecular Weight	392	380
ASTM Ash, wt-%	0.22	0.50
Con. Carbon, wt-%	9.43	12.3
C ₅ Insol., wt-%	21.3	23.5
C7 Insol., wt-%	19.1	22.8
Toluene Insol., wt-%	6.12	7.12
Bromine Number	6.6 (Modified	
	Method)	
Carbon, wt-%	84.44	84.85
Hydrogen, wt-%	12.45	12.77
Sulfur, wt-%	1.25	1.70
Nitrogen, wt-%	0.52	0.524
Oxygen, wt-%	1.64	0.88 ± 0.06
Emission, wt-ppm		
Fe	780	54
N i	9	4.8
V	86	13
Ca	180	6.9
Mg	0.11	1.4
Mn	25	1.6
Cr	5.0	0.51
Sn	33	0.99
Cu	61	2.5
Zn	100	10.6
Ti	9.7	6.7
Pb	7.5	0.78
Na	78	9.2
Мо	8.9	0.85
Co	2.2	_
Si	23 0	Diluent
A1	Diluent	3.4
В	50	0.78
Ва	1.7	1.6
Sr	••	-
Zr	0.36	-
Ash	2,780	709
Melting Point, °F		206*

^{*} Heavy material left on tip of thermometer which will not melt.

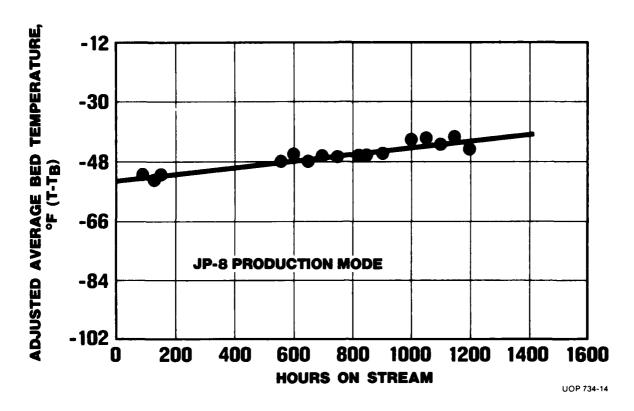


FIGURE 5

PARALLEL — FLOW HYDROCRACKING
CATALYST ACTIVITY AND STABILITY

SECTION IV

PLATFORMING OF A C7-300°F SHALE OIL NAPHTHA

As part of the Phase III program, a narrow boiling naphtha cut from the hydrocracking operations was catalytically reformed to obtain basic information on yields and octane, and to compare this shale-derived naphtha with petroleum stocks.

Shale oils generally have narrower boiling ranges, from about 400° to 1000°F, and contain less naphtha as compared to petroleum crudes. With this boiling range, shale oils readily lend themselves to production of diesel and jet fuels. However, high yields of good quality gasoline can be produced from shale oil depending on the refinery configuration.

The UOP Platforming® process is a proprietary catalytic reforming process developed by UOP. It can be used to produce an excellent petrochemical feed for BTX recovery, or produce a high octane blending component for gasoline production. A valuable by-product of these operations is the production of hydrogen. In this instance, Platforming was used to investigate the shale oil naphtha as a potential source of special motor fuel.

Pilot Plant Description

A schematic diagram of the pilot plant used for this operation is shown in Figure 6. Fresh feed with recycle hydrogen is fed over the catalyst. The reactor effluent is condensed and the gas and liquid separated. The gas is recycled back to the reactor while the propane and heavier liquid is debutanized. The hydrogen produced in the process is removed continuously through a pressure regulator, measured and analyzed. The debutanized C5+ product is similarly weighed and analyzed.

Present commercial Platforming catalysts comprise extruded or spherical alumina promoted with Group IV, VII and VIII metals. A commercially proven UOP Platforming catalyst designated Rx-320 was employed in this study.

Feedstock

The Platforming naphtha feed was prepared by fractionating the hydrocracked product from the Phase II operations in two steps to remove the 300° F+ heavies and the C_6 - light front end to yield essentially 10 gallons of a C_7 to 300° F product. Inspections of this product are shown in Table 22.

This 58.5 °API gravity naphtha contained about 91 vol-% paraffins plus naphthenes, 1 vol-% olefins and 8 vol-% aromatics. The clear Research octane number was about 55. The nitrogen content was acceptable at <0.1 ppm, but the 110 ppm sulfur content and the 1% olefins were outside the normal specifications for Platformer feedstock. The high sulfur content was a result of the sulfur addition experiments in the hydrocracking program.

A mild hydrotreating operation was used to prepare the Platformer[®] feed. The operating conditions were typical of those required to hydrotreat a petroleum-derived naphtha. Following this hydrotreating step, the total product was fractionated to reduce the 95% and end points to about 255° and 298°F, respectively. The analysis of the Platformer feed is shown in Table 23. The sulfur and nitrogen contents are well below the levels required for current Platforming operations. The aromatics content has been reduced to about 5 vol-%. A detailed GC analysis of this material is given in Table 24.

Operating Conditions

Process conditions used in this study are comparable to those commercially employed for reforming petroleum naphtha. The base conditions selected for reference represent those employed commercially for UOP's "State-of-the-Art" design of Continuous Platforming units. The nomenclature adopted to relate the actual operating conditions to these base conditions are as follows:

		Written as
Temperature	T-T (base), °C	τ - τ_B
Pressure	P-P (base), psi	P-PB
Space Velocity	LHSV/LHSV (base)	L/LB
Hydrogen/Feed Ratio	H ₂ /Feed	H/HB
, , , , , , , , , , , , , , , , , , , ,	H ₂ /Feed (base)	

This operation covered a range of temperatures from 70°F below to 20°F above the base temperature to obtain a yield-octane relation for this feed. The other operating conditions were:

Pressure	P-PB, psi	0	
Space Velocity	L/LB	2	
Hydrogen/Feed Ratio	н/н _в	2.5	

Yield-Octane Study

The Platformer product analyses with Research Octane numbers from 90 to 104 are given in Table 25. Product aromatic contents range from about 50 vol-% at 90 RONC to about 79 vol-% at 104 RONC. The product octanes responded very well to lead addition. There is an 8.5 octane increase in the product from the lowest octane operation to a 5.0 increase at the highest octane.

The product yield data are related to octane level in Figures 7, 8 and 9. The C₅ plus liquid yield varied linearly from 85 vol-% at 89 RONC to 70 vol-% at 104 RONC. At the highest operating severity there is an indication that the hydrogen yield (Figure 8) is leveling off.

Overall, the data show this shale oil naphtha cut to be an excellent Platformer feedstock that will give yields comparable to petroleum derived feeds when reformed to octanes ranging from 90 to 104 RONC.

Special Gasoline Preparation

The individual products from this Platforming operation were blended to make a larger sample which was subsequently used to make a Special Grade, Class B volatility, gasoline. The analyses of this sample, as reported in Table 26, show a 47.2 °API gravity material having a Research octane number of 95.5.

To meet all the Class B volatility specifications, calculations showed it would be necessary to blend together 60 vol-% of the Platformate® blend, 30 vol-% of a C_5 - C_6 cut previously distilled out from the hydrocracked shale oil naphtha product, and 10 vol-% of butanes. For practical considerations in the laboratory, the 10 vol-% of butanes was not blended with the liquids noted above.

An analysis of the C_5 - C_6 fraction having a Research octane number of 70.5 is given in Table 27.

The actual gasoline blend sent to the USAF consisted of 2:1 volume blend of Platformate and the C5-C6 cut. Analyses of this blended gasoline are given in Table 28 and show that this gasoline meets all the chemical and antiknock requirements as outlined in Federal Specification VVG-1690B, July 1, 1978. A clear Research octane number of 87.2 was determined for this blend. A detailed isomer distribution analysis is given in Table 29.

The final Special Grade gasoline blend calculated by adding the equivalent of 10 vol-% butanes to the blended gasoline is shown in Table 30. This blend does not meet the Class B volatility specs for vapor pressure and TVL at 20°C but falls between the Class B and C gasolines. A slight adjustment in butane concentration would be sufficient to bring the gasoline either to a Class C or B volatility.

Conclusions

Naphtha derived from hydrocracking shale oil can be catalytically reformed to yield high octane gasoline blending components. The C5 plus

liquid yield varied linearly from 85 vol-% at 89 RONC to 70 vol-% at 104 RONC. Overall, this shale oil naphtha cut proved to be an excellent Platformer feed and gave yields comparable to petroleum-derived naphtha.

TABLE 22. NAPHTHA SAMPLE FROM HYDROCRACKING

Analysis		GLC	<u>₩t-%</u>
API Gravity at 60°F	58.5	<u>i</u> -C ₄	-
Sp.Gr. at 60°F	0.7447	\overline{n} -C ₄	-
Distillation (D-86), °	F	<u>1</u> -C5	_
IBP	217	<u>n</u> -C ₅	-
5%	224	<u>2</u> −MP	-
10%	227	3-MP	-
20%	230	2,2 DMB	_
30%	234	2,3 DMB	_
40%	240	$\underline{\mathbf{n}}$ - \mathbf{c}_6	_
50%	246	_C ₇ ₽	16.72
60%	250	C ₈ P	22.00
70%	256	C ₉ P	14.49
80%	266	c_{10} P	2.45
90%	280	c_{11}^{10} P	
95%	293	11	
EP	338	C-C5	_
% Over	99.0	MCP	-
% Botts.	1.0	c-c ₆	0.02
% Bocco	• • •	C ₇ n	11.96
Sulfur, ppm	110.5	C ₈ n	13.90
Nitrogen, ppm	<0.1	Cgn	8.15
Oxygen, ppm	117	C ₁₀ n	0.91
oxygen, ppm	• • •	C ₁₁ n	
Carbon, wt-%	82.01	-11	
Hydrogen, wt-%	14.35		
Mercaptan, ppm	0.5	Benzene	0.0
FIA, vol-%	0.0	benzene	0.0
A A	7.8	Toluene	2.9
Ô	1.1	C _R arom	4.8
P + N	91.1	Cg+ arom	0.7
r + N	71.1	Total	8.4
		10242	3 • •
RON, Clear (micro)	54.6		
		Olefins	1.0

TABLE 23. HYDROTREATED PLATFORMER FEEDSTOCK

API Gravity at 60°F	62.0
Sp. Gr. at 60°F	0.7313
Distillation, (D-86), °F	
IBP	194
5%	204
10%	208
20%	214
30%	216
40%	220
50%	224
60%	227
70%	232
80%	238
90%	248
95%	254
EP	298
% Over	99. 0
Molecular Weight	104
Sulfur, ppm	<0.1
Nitrogen, ppm	0.20
Carbon, wt-%	83.38
Hydrogen, wt-%	15.04
Chloride, ppm	4.4
Oxygen, ppm	< 10
Mercaptan Sulfur, wt-%	0.0001
Lead, ppb	< 20
Arsenic, ppb	1.4
Bromine Index	131.7
RON, Clear (micro)	56.4
PONA, vol-%	
P	61
0	0
N	34
A	5
FIA, vol-%	
A	5.5
0	-
P + N	94.5

TABLE 24. COMPONENT ANALYSIS OF HYDROTREATED PLATFORMER FEEDSTOCKS

GC Analysis, Wt-%

Aromatics		ECP	2.59
Toluene	3.8	2,5-DMH	1.42
EB	0.5	2,4-DMH	1.48
p-Xylene	0.5	1-t-2-C-4-t-MCP	1.73
m-Xylene	1.1	1-t-2-C-3-t-MCP	0.92
o-Xylene	0.4	3,3-DMH	0.21
Total Aromatics	6.3	1,1,2-TMCP	0.41
		2-M-3-DP + 2,3-DMH	1.07
Paraffins and Naphthe	nes	1-c-2-t-4-t-MCP	0.53
<u>i-C5</u>	0.44	1-c-2-t-3-t-MCP	0.34
n-C5	0.39	?-MH + 3-M-3-EP	5.94
$\overline{2}$, 2-DMB	0.02	1-c-2-c-4-5-MCP + 4-MH + 3,4-DMH	3.40
Cyclo C ₅	0.02	3-EH + 3-MH + 1,1-DMCH	6.68
2,3-DMB	0.17	1-t-4-DMCH + 1-C-3-DMH	2.28
2-MP	1.27	1-MC-3-ECP	2.02
3-MP	0.85	1-M-t-3-2-ECP + 1-M-1-ECP	2.90
n-C ₆	1.22	1-C-2-C-3-t-MCP + 1-t-2-DMCH	1.04
MCP	0.95	$1-t-3-DMCH + 1-c-4-D\overline{M}CH$	1.89
2,2-DMP	0.05	IPCP —	0.17
2,4-DMP	0.19	n-0	4.71
Cyclo C ₆	0.21	1-M-c-2-ECP	0.54
3,3-DMP	0.09	$1-c-\overline{2}-DMCH$	0.46
1,1-DMCP	0.44	ECH + n-P-c-P	1.45
2-MH	6.65	Co Naphthenes	2.23
2,3-DMP	1.58	Cg Paraffins	3.71
1-cis-3-DMCP	2.61		
3- MH	7.03	Total $P + N$	93.7
1-t-3-DMCP	1.91		
$1-\overline{t}-2-DMCP$	2.14		
3-EP	0.57		
n-C ₇	6.76		
1-C-2-DMCP	1.48		
мсн	4.77		
1,1,3-TMCP	0.94		
2,2-DMH	0.23		

See Table 29 for compound definitions.

TABLE 25. PLATFORMER PRODUCT ANALYSES

Period	1	2	3	4	5	6	7	8	9
API Gravity at 60°F	_		47.8			44.3		40.2	50.9
Sp. Gr. 60/60°F 0									
Distillation (D-86),							-,		
IBP	129	144	136	136	128	122	120	121	140
5%	175	178	166	165	158	154	154	158	178
10%	188	190	182	180	176	173	177	178	191
20%	206	206	201	200	200	199	208	209	206
30%	217	217	216	215	217	218	230	232	218
40%	226	225	226	227	230	232	242	244	226
50%	234	234	236	238	241	243	250	252	233
60%	242	242	246	252	251	252	256	258	240
70%	251	253	256	259	260	260	264	265	250
80%	264	265	268	268	270	270	273	274	262
90%	282	282	284	283	285	283	286	285	280
95%	296	295	300	296	300	299	300	300	298
EP	350	328	331	337	336	338	346	356	316
% Over	99.0	99.0	99.0	99.0	99.0	99.0	99. 0	99.0	99. 0
% Bottoms	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
FIA, vol-%									
A	50.2	52.4	57.2	58.9	64.8	67.1	77.7	79.2	49.1
0	1.3	1.3	1.3	1.3	1.3	1.4	1.1	.9	1.6
P + N	48.5	46.3	41.5	39.8	33.9	31.5	21.2	19.9	49.3
Octane Number									
RONC	90.2	91.2	95.2	95.9	98.2	99.8	103.7	104.3	88.7
RON + 3 TEL	98.7	99.3	102.3	102.0	104.2	104.7	107.6	109.2	97.6
MON	80.6	81.2	84.4	86.8	87.3	88.5	92.0	92.4	79.5

TABLE 26. PLATFORMER PRODUCT BLEND

API Gravity at 60°F	47.2
Sp. Gr. 60/60°F	0.7918
Distillation (D-86), °F	
IBP	137
5%	175
10%	189
20%	207
30%	221
40%	230
50%	238
60%	247
70%	257
80%	270
90%	287
95%	306
EP	351
% Over	97.5
% Bottoms	1.0
% Loss	1.5
Dry Point, °F	320
RON Clear	95.5
MON Clear	84.8
Reid Vapor Pressure, psi	3.3

TABLE 27. C₅-C₆ FRACTION

API Gravity at 60°F	77.3	GC	<u>₩t-%</u>
Sp. Gr. 60/60	0.6777	<u>1</u> -C ₄	trace
Distillation (D-216),	°F	\underline{n} -C ₄	0.1
IBP	120	<u>i</u> -C ₅	7.6
5%	130	<u>n</u> -C ₅	8.7
10%	134	2,2-DMB	0.3
20%	137	CP	0.6
30%	140	2,3-DMB	2.7
40%	144	2-MP	19.5
50%	148	3-MP	13.0
60%	152	\underline{n}^{-C}_6	16.9
70%	155	MCP	13.1
80%	159	СН	1.9
90%	166	Bz	0.9
95%	174	C ₇ +	14.6
EP	240	Olefins	0.1
% Over	99. 0		
% Bottoms	1.0		
Nitrogen, ppm	0.13		
Sulfur, ppm	32		
Vapor Press., psi	7.7		
RON Clear	70.5		
MON Clear	68.8		

TABLE 28. GASOLINE BLEND

API Gravity at 60°F Sp. Gr. 60/60°F Distillation (D-86), °F	56.2 0.7539	Existent Gum, mg/100 mL Unwashed Gum, mg/100 mL Oxidation Stability, min. >12	1 I 200
IBP	151	RON Clear 87	7.2
5%	151		9.8
10%	170		3.5
20%	180	KTM/ Z	,,,
30%	192	Water and	
40%	206	Sediment, vol-% <0.0	105
50%	208	Sediment, Voi-%	JOJ
60%		Panana19	1.2
70%	244	- · · · · · · · · · · · · · · · · · · ·	1 • 2
80%	261	Phosphorus (unleaded), g/gal <0.00	003
90%	282	g/gal <0.00	JU 3
95%	306		
EP	323		
% Over	98.0		
% Loss	1.0		
% Bottoms	1.0		
Dry Point, °F			
Carbon, wt-%	85.86		
Hydrogen, wt-%	13.70		
Nitrogen, ppm	0.3		
Oxygen, ppm	352		
Sulfur			
(unleaded), ppm	5.6		
Reid Vapor Pres., psi	5.1	(35.16 kPa)	
FIA, vol-%			
A	39.0		
0	1.1		
P + N	59.9		
Cu Strip Corrosion	1 A		
Autiknock Content			
Lead (unleaded)	< 0.01	g/gal	
Manganese (unleaded)	< 0 • 1	ppm	
Temperature, °C, min. at V/L Ratio = 20	71.3		
	"		

TABLE 29. GASOLINE BLEND

Complete Isomer Distribution

	Wt − %
Aromatics	
Alonacics	
n	1 5
Benzene	1.5
Toluene	18.5
Ethylbenzene	2.7
<u>p</u> -Xylene	4.7
<u>m</u> -Xylene	9.6
o-Xylene	5.9
C ₉ + Aromatics	4.2
Total A	46.9
Olefins (Total)	1.0
Paraffins and Naphthenes	
1. Propane	
2. Isobutane	
3. n-Butane	0.10
4. Isopentane	4.49
n-Pentane	4.04
**	0.14
<pre>. // 2-bimethylbutane Cvclopentane</pre>	0.23
	0.56
6. 2,3-Directhylbutane 9. 3-Merbylpentane	
· · · · · · · · · · · · · · · · · · ·	4.30
10. (-Methylpentane	3.22
11. <u>n</u> =tiexane	5.18
12. Methylavelopentane	3.03
13. 2,1-Dimethylpentane	0.59
14. 2,4-i)fmethylpentane	1.19
15. 2,2,3-Trimethylbutane	0.10
16. Cyclohexane	0.54
17. 3,3-Nimethylpentane	0.47
18. 1,1- methylcyclopentane	0.11
19. 2-Yethylhexane	3.93
20. 2, -Dinethylpentane	1.28
21. 1-c :-3-Dimethylcyclopentane	0.10
22. 3-Mernylhexane	4.50
23. 1-trans-3-Dimethylcyclopentane	0.15
24 1-trans-2-Dimethlycyclopentane	0.16
25. 3-Ethylpentane	0.44
26. 2,2,4-Trimethylpentane	0.02
27. n-Heptane	3.50
28. T-cis-2-Dimethylovolopentane	0.02
	0.03
19. Met by Laye Lohexane	0.02
30. 1,1,3-Trimethylovolopentane	0.25
11. ','-'Imerlablicane	
2. st. wisy lopertane	0.04
33. 1, -Dinethylbexane	0.38
3 ,4=1 (me*) withexame	0.70
3. , , selrim t vipentane	0.01
36. Terrans-Legis-4-Trimethyleyelop	entane

A

TABLE 29. GASOLINE BLEND (Continued)

Paraffins and Naphthenes (Cont.)

37.	l-trans-2-cis-3-Trimethylcyclopentane	0.01
38.	3,3-Dimethylhexane	0.27
39.	2,3,4-Trimethylpentane	0.03
40.	1,1,2-Trimethylcyclopentane	0.01
41.	2,3,3-Trimethylpentane	0.03
42.	2-Methyl-3-ethylpentane	0.51
43.	2,3-Dimethylhexane	0.51
44.	l-cis-2-trans-4-Trimethylcyclopentane	
45.	1-cis-2-trans-3-Trimethylcyclopentane	
46.	2-Methylheptane	1.52
47.	3-Methyl-3-ethylpentane	1.02
48.	l-cis-2-cis-4-Trimethylcyclopentane	
49.	4-Methylheptane	1.07
50.	3,4-Dimethylhexane	
51.	3-Ethylhexane	
52.	3-Methylheptane	2.35
53.	l,l-Dimethylcyclohexane	
54.	l-trans-4-Dimethylcyclohexane	
55.	1-cis-3-Dimethylcyclohexane	
56.	1-Methyl-cis-3-ethylcyclopentane	0.03
57.	1-Methyl- <u>trans</u> -3-ethylcyclopentane	
58.	l-Methyl-trans-2-ethylcyclopentane	0.07
59.	l-Methyl-l-ethylcyclopentane	
60.	l-cis-2-cis-3-Trimethylcyclopentane	
61.	l-trans-2-Dimethylcyclohexane	
62.	I-trans-3-Dimethylcyclohexane	
63.	l-cis-4-Dimethylcyclohexane	
64.	Isopropylcyclopentane	
65.	n-Octane	1.71
66.	1-Methyl-cis-2-ethylcyclopentane	
67.	l-cis-2-Dimethylcyclohexane	0.04
68.	Ethylcyclohexane	0.02
69.	<u>n</u> -Propylcyclopentane	0.02
	C ₉ Naphthenes	0.09
	Cg Paraffins	0.39
	C ₁₀ Naphthenes	0.13
	C ₁₀ Paraffins	
	C ₁₁ Naphthenes	
	C ₁₁ Paraffins	
	$C_{12}^{11} P + N$	
	Total P + N:	52.1

Totals

TABLE 30. SPECIAL GASOLINE

API Gravity at 60°F	60.5
Sp. Gr. 60/60°F	0.7370
Distillation (D-86), °F	°F
IBP	96
10%	125
30%	168
50%	195
70%	235
90%	278
EP	306
RON Clear	87.9
MON Clear	80.8
$\frac{R + M}{2}$	84.4
	psi
Reid Vapor Pressure	11.1
Temperature, °C min at V/L Ratio = 20	53.0

d

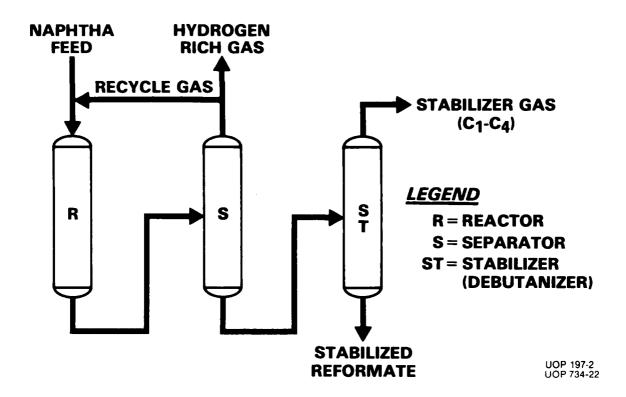
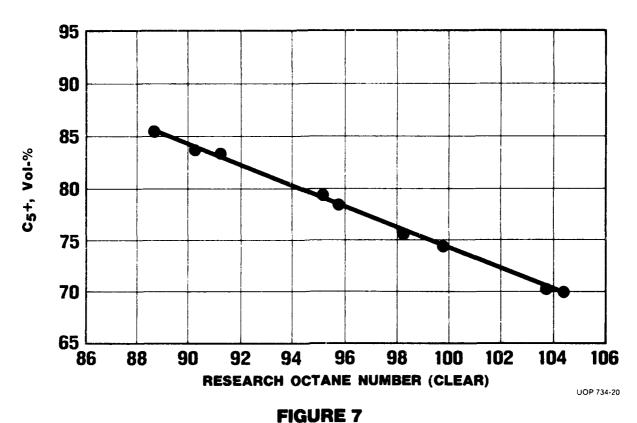
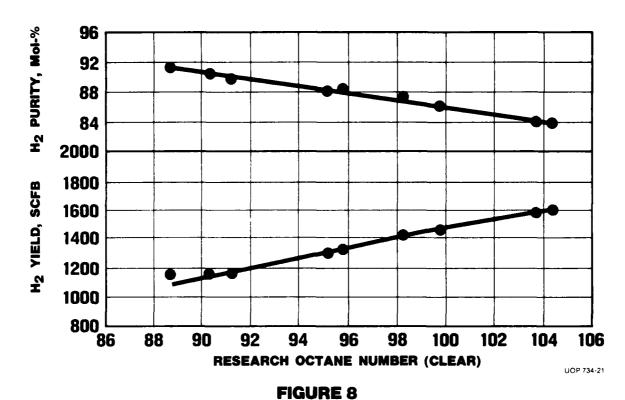


FIGURE 6
NAPHTHA REFORMING PILOT
PLANT

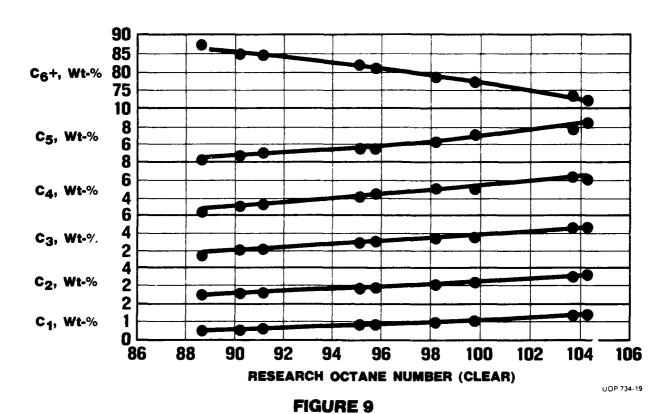
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PLATFORMING YIELD — OCTANE STUDY
OCCIDENTAL SHALE OIL
LIQUID AND HYDROGEN YIELDS



PLATFORMING YIELD — OCTANE STUDY OCCIDENTAL SHALE OIL LIQUID AND HYDROGEN YIELDS



PLATFORMING YIELD — OCTANE STUDY OCCIDENTAL SHALE OIL

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SECTION V

ARSENIC MANAGEMENT STUDIES

Shale oils produced by current retorting operations contain arsenic in such concentrations that it deserves special attention. Two approaches to the shale oil arsenic management problem have been investigated in this program:

- 1. Crude shale oil arsenic solubilization
- 2. Deposited arsenic passivation or extraction.

Shale Oil Arsenic Solubilization

The first hypothesis investigated involved converting the arsenic in the raw shale oil to a water soluble compound. This might be accomplished by injecting a reagent downstream of the retort into the oil-water mixture. With intimate mixing of the reagent, oil and water may convert the arsenic into water soluble compounds. After the two phases are allowed to coalesce, the water phase is drained off leaving a shale oil with reduced arsenic content. The water containing arsenic could then be further treated, if necessary, to render it environmentally safe.

Desalted Occidental shale oil with 19 ppm arsenic was utilized as the hydrocarbon source for a series of experiments attempting to convert arsenic into water soluble compounds. For these experiments, the water, reagent and shale oil were intimately contacted utilizing a shear-type mixer. Each test was performed by mixing the oil and water for 15 minutes at about 95° C (200°F) . The type of emulsion formed while adding various reagents was observed, and the arsenic level left in the oil was measured after each test.

A summary of results is shown in Table 31. Arsenic solubilization ranged from 13 to 52%. The removal of arsenic was probably accomplished by

the formation of an insoluble material that is associated with the emulsion. The use of a wide range of reagents along with shear mixing to effect water solubilization of arsenic from shale oil does not appear to be very promising.

Arsenic Passivation and Extraction

The high concentration of arsenic in the raw shale oil and its removal and containment on the first-stage hydrotreater catalyst poses some special handling problems. Before arsenic-laden catalysts can be disposed of as non-hazardous wastes, the aqueous solubility of arsenic as measured by the EPA Toxicity test must be reduced to less than 5 ppm. In an effort to meet this requirement for safe disposal, the following objectives were considered:

- Ascertain the composition of arsenic-containing species on used catalyst.
- 2. Determine the arsenic solubility of aqueous extracts of used catalysts.
- 3. Evaluate methods of passivating soluble arsenic.
- 4. Determine the effect of various gases on arsenic volatility.
- 5. Evaluate the extraction behavior of arsenic from untreated and thermally treated used catalysts using various solvents.
- 6. Determine the conditions for complete dissolution of used catalysts.

Experimental Procedure

Samples of spent catalysts were thermally treated by spreading the catalyst pills inside a 1-inch silica tube. Three thermocouples were placed at the ends and middle of the catalyst bed. The catalyst sample was held in

place by tufts of quartz wool. Rubber stoppers fitted with glass tubes allowed controlled atmospheres to be passed through the sample gas flow as adjusted to 600 mL/min.

Heating was accomplished by a 30 cm-long split tube furnace. Once the desired temperature had been reached, the silica tube containing the sample was inserted. The furnace temperature was held to within $\pm 10^{\circ}$ C. Cooling was achieved by removing the apparatus from the furnace. When cool, the catalyst was removed from the tube. The sample was analyzed for arsenic and sulfur.

Thermally treated and as-received samples were extracted in various solvents. Five gram samples were added to Erlenmeyer flasks containing 200 mL of extractant and shaken on a wrist shaker for 3 hours. The resulting slurry was filtered and washed. The catalyst residue and filtrate were analyzed for arsenic and sulfur.

Elemental Analysis

The elemental analyses of spent catalyst beds removed from two first-stage hydrotreating pilot plant reactors show that the arsenic concentration declines rapidly downward through the bed. As shown in Table 32, iron exhibits a similar profile while the remaining elements are fairly evenly distributed. The major difference between the two catalysts is the much higher arsenic level on the used catalyst designated as "Catalyst 2". The relative concentrations of the arsenic and iron throughout the catalyst bed are not significantly different.

An identification of the arsenic compounds present on the catalyst was considered necessary in order to determine the best methods to either fix the leachable arsenic by further chemical reaction or convert the arsenic into a highly soluble form. X-ray diffraction patterns obtained for the two samples were analyzed and the results are summarized in Table 33. The peak positions were calculated, and a manual search was made of the powder diffraction files in order to determine which compounds were present.

The Υ -Al₂O₃ of the catalyst base is a major component in both samples. Trace levels of α -quartz and boehmite were also detected. The rest of the standard patterns listed fit the sample patterns well, but the actual components present may have slightly different compositions.

Both samples contain a major phase similar to pyrrhotite $[Fe_{(1-x)}S$, where $x \le 0.1]$. A slight increase in sulfur content would lead to a mixture of pyrrhotite and pyrite (FeS_2) . There is evidence of pyrite in the sample designated "Catalyst 1".

 ${\rm Co_{0.84}Ni_{0.16}As_{1.04}}$ is the cobalt analog of niccolite (NiAs)₁. Their patterns are similar. The peaks in the pattern more closely match the former (${\rm Co_{0.84}Ni_{0.16}As_{1.04}}$); however, the actual compound present may have a formula somewhere between the two arsenides.

Although some of the peaks for the two used catalysts match the pattern of $\text{Cu}_{24}\text{As}_{12}\text{S}_{31}$, there was no copper detected in the elemental analysis. It is possible a similar compound was present with other metals substituted for the copper.

The chloroform-washed samples were subjected to a modified EPA Toxicity test for arsenic. The modification in procedure was the use of $10~\rm grams$ in place of $i00~\rm gram$ samples. As shown in Figure 10, the As solubility of the chloroform-washed catalyst appears to be a linear function of the As content of the catalyst. In order to meet the EPA limit, the catalyst would have to contain less than $0.2~\rm wt-\%$ As.

Metal salts were blended with the used catalyst, then heated in an inert atmosphere in an attempt to fix the arsenic as metal arsenides or metal arsenous sulfides. It was anticipated that conversion to these compounds would cause the arsenic to be less soluble; however, as shown in Figure 10, this proved not to be the case.

The EPA test was performed on several used catalysts that had been previously extracted with sulfuric and nitric acid solutions. The arsenic solubility was decreased; however, the arsenic level remaining on the

catalyst was still too great for direct disposal. As shown in Figure 10, in order to meet the EPA limit, it will be necessary to reduce the As content on the catalyst to about 0.5 wt-%.

If essentially all of the arsenic could be volatilized in a roasting process, arsenic collection would be feasible. Considering the relatively high vapor pressures of arsenic, arsenic sulfide and arsenic oxide, it appears that arsenic would be removed at elevated temperatures in either oxidizing, reducing, or neutral atmospheres.

As shown in Figure 11, the volatility of arsenic from spent catalyst was dependent upon temperature, residence time and atmosphere. It appears that a temperature of about 500°C (932°F) is required to break down the original arsenic minerals and to ensure adequate vapor pressure of volatile species. If the rate of arsenic volatilization can be increased, arsenic might be recovered by such a process.

Extractants were chosen to discern the state of arsenic present on the catalyst following thermal pretreatment. $H_2SO_4/FeCl_3$ and HNO_3 are strong oxidizing agents capable of oxidizing sulfide minerals. Sulfuric acid is a non-oxidizing acid which can solubilize sulfates and arsenates. Sodium sulfide forms complex anionic arsenous sulfides which are highly soluble.

Arsenic extraction results, from catalysts that were thermally oxidized at the severe conditions of 500° C in 5% 0_2 , were very similar. The data shown in Figure 12 indicate that the arsenic has probably been converted to an arsenate.

The data plotted in Figure 13 indicate that catalysts pretreated under rather neutral conditions, 25% H₂0/75% N₂, were more effectively extracted with oxidizing acids than with the other leachants. The arsenous sulfide minerals are only beginning to be broken down at the highest temperature, 600°C (1112°F).

As shown in Figure 14, using SO_2 in the thermal treatment step produces a different response to the extractants. At temperatures below 500° C, the

 ${\rm SO}_2$ atmosphere produced little change in the arsenic chemistry. At $500^{\circ}{\rm C}$, ${\rm SO}_2$ reacted slowly with carbon to form sulfur and ${\rm CO}_2$ and converted the metal arsenous sulfides to more leachable forms. All of the extractants remove approximately the same level of arsenic.

None of the thermal pretreatments produced a material that allowed high levels of arsenic extraction. In the best case, about 80% of the arsenic was removed by a combination of thermal treatment and leach extraction. Increasing the leachant concentration to increase the arsenic extraction would result in considerable alumina dissolution.

An alternative approach to arsenic recovery utilized solution oxidation. Acid digestion of as-received used catalyst was performed in a stoichiometric quantity of sulfuric acid, assuming the catalyst composition to be entirely Al_2O_3 with the following reaction:

$$Al_2O_3 + 3 H_2SO_4 \rightarrow Al_2(SO_4)_3 + 3 H_2O$$

The digestion was carried out under reflux for 8 hours. In order to obtain high recoveries of the elements, the reaction had to be carried out at a positive solution potential. This was accomplished by adding 3 wt-% HNO3 to the sulfuric acid. As shown in Table 34, 90% recovery of arsenic was achieved with both the 10 and 20 wt-% sulfuric acid solutions. Only molybdenum was poorly recovered in this solvent; however, molybdenum can be recovered from the residue using other techniques. Digestion of thermally treated catalysts was not as effective in recovering arsenic as solution exidation of the as-received used catalyst.

Conclusions

- Used catalysts from processing shale oil will contain higher levels of soluble arsenic than allowed by the EPA for non-hazardous disposal.
- 2. Much of the arsenic is present on the catalyst as thermally stable metal arsenides and metal arsenous sulfides.

- 3. It was not possible to fix or passivate the arsenic on the catalyst in an insoluble form by thermally treating the catalyst with or without additives.
- 4. Leachants, which do not dissolve excessive quantities of alumina, extract only about 20% of the arsenic from non-thermally treated catalysts.
- 5. Of the thermal treatments investigated, only dilute oxidizing atmospheres, 5% O_2 or less, at elevated temperatures, $\geq 500^{\circ}\text{C}$, for extended time result in appreciable arsenic volatilization. This indicates that arsenic is not present on the catalyst as a simple sulfide, oxide or metallic compound.
- 6. Increasingly severe thermal pretreatments allow up to 75% of the arsenic to be extracted by dilute leachants.
- 7. Both temperature and gas composition are important variables in converting the arsenic to a leachable form.
- 8. Results of toxicity tests on previously treated catalysts indicate that high extraction levels are necessary before the residue is acceptable.
- 9. Digestion of as-received spent catalyst requires a high solution potential in order to achieve good extractions.
- 10. Digestion of thermally oxidized catalysts yields poorer arsenic extractions than solution oxidation of the as-received catalysts.
- 11. Topics worthy of further investigation include the evaluation of different ratios of As, Fe, C and S on arsenic volatility and solubility; the recovery of arsenic by thermal means; the development of methods to recover arsenic from acidic solutions; and the determination of the minimum acid necessary to achieve good extractions of arrenic without excessive dissolution.

TABLE 31. SHALE OIL ARSENIC SOLUBILIZATION STUDY

Arsenic in Feed = 19 ppm

Reagent	Observation	Arsenic in Oil, ppm
10% кон	Emulsion stable, broken by addition of isooctane/methanol.	10.8
10% H ₂ SO ₄	No emulsion formed.	16.6
50% Acetic acid	Formed emulsion which slowly broke.	14.7
10% Na ₂ S	Formed emulsion which did not break with isooctane/methanol treatment.	
	1) One hour at 220°C (430°F) under 100 ATM of N ₂ -broke.	11.7
	2) Centrifuged at 7000 ppm, part of the emulsion broke.	9.1

TABLE 32. FIRST-STAGE HYDROTREATER, SPENT CATALYST ANALYSIS

Sample Location			E1	ement,	wt-%			
Catalyst 1	<u>Fe</u>	<u>C</u> _	N	<u> </u>	<u>H</u>	_As_	Mo	Co
Upper Section	6.3	10.4	0.47	9.3	1.6	2.4	+	+
Upper Middle	4.5	10.5	0.34	8.6	1.4	1.2	+	+
Middle Section	1.0	13.4	0.57	8.2	1.6	0.32	+	+
Lower Middle	0.4	13.3	0.59	7.8	1.4	0.12	+	+
Lower Section	0.4	14.1	0.67	7.5	1.3	0.07	+	+
Catalyst 2								
Тор	6.0	9.11	3.9	6.2		8.4	+	+
Top Middle	3.7	10.2	0.43	6.4		5.6	+	+
Middle	1.9	10.1	0.43	5.7		3.4	+	+
Bottom Middle	1.0	10.2	0.85	5.8		1.6	+	+
Bottom	0.9	10.3	0.22	5.5		1.3	+	+

+ = Present.

TABLE 33. DIFFRACTION DATA ON USED CATALYSTS

Compound	Catalyst l	Catalyst 2
Identification	Upper Section	Upper Section
YA1 ₂ 0 ₃	m	m
asio ₂	t	t
AlO(OH) Boehmite	t	
Fe(1-x)S Pyrrhotite	m	m
FeS ₂ Pyrite		t
Co _{0.84} Ni _{0.16} As _{1.04}	t	m
Cu ₂₄ As ₁₃ S ₃₁	t	m

m = major

t = trace

TABLE 34. DIGESTION OF USED CATALYST

Used Catalyst Analysis: Arsenic, wt-% 1.16

Digestion	Solution	As Extraction, %
H_2SO_4 , wt-%	$\frac{\text{HNO}_3}{\text{MNO}_3}$, wt-%	, , , , , , , , , , , , , , , , , , ,
10	-	24
10	3	90
20	-	23
20	3	90
40	-	24
40	3	72

Thermally Treated Catalyst Extracted in 40 wt-% $\rm H_2SO_4$

Therma	al Treat	nent	%	As Removed	
Temp., °C	Gas	Time, Hr	Volatilization	Extraction	Total Removal
5 50	5% 0 ₂	6	69	50	74
650	5% 0 ₂	2	58	53	80
250	5% 0 ₂	6	9	74	76

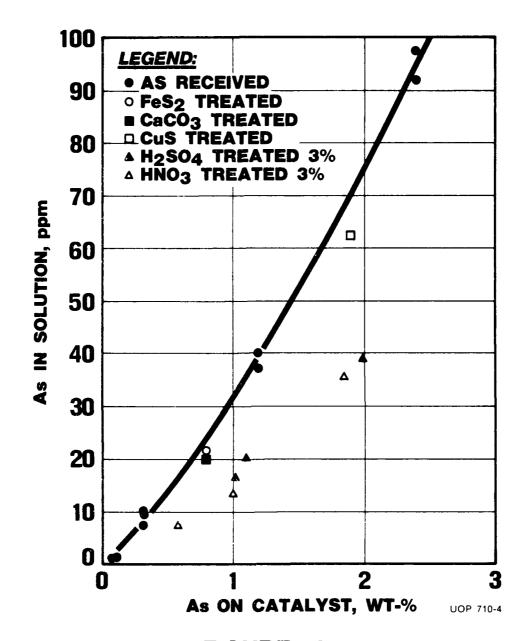


FIGURE 10
SIMULATED EPA TOXICITY
TEST ON USED CATALYST

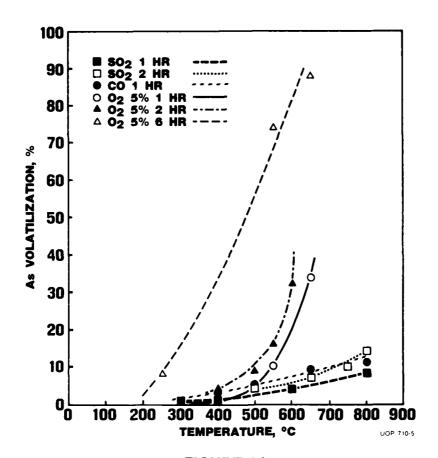


FIGURE 11
EFFECT OF THERMAL TREATMENT
ON ARSENIC VOLATILIZATION

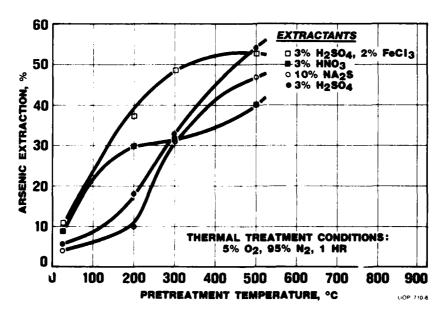


FIGURE 12
EFFECT OF SEVERE OXIDATION
ON ARSENIC EXTRACTION FROM
USED CATALYST

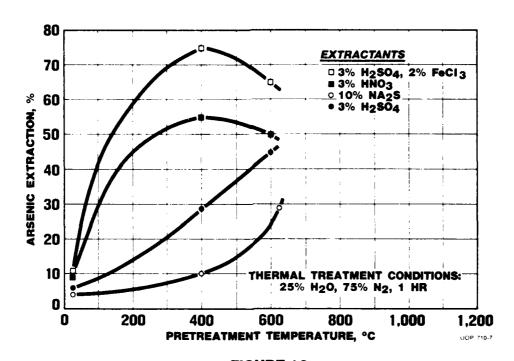


FIGURE 13
EFFECT OF MILD OXIDATION ON ARSENIC EXTRACTION FROM USED CATALYST

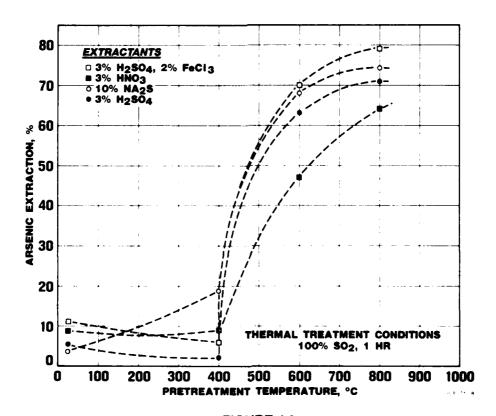


FIGURE 14
EFFECT OF SULFUR TREATMENT ON ARSENIC
EXTRACTION FROM USED CATALYST

SECTION VI

FOULING

The Monirex® Fouling Monitor has been used to determine the thermal fouling properties of aged Occidental desalted shale oil, an Arabian Light Berri petroleum oil and blends of these oils. In addition, the thermal fouling properties of a severly hydrotreated Occidental shale oil, a JP-8 fuel derived from the hydrotreated shale oil and a relatively fresh Paraho shale oil have also been determined.

Experimental Development

The UOP Monirex Fouling Monitor, previously described in considerable detail in the Phase II report, was used to evaluate the relative thermal fouling properties of shale and petroleum oils. This unit measured the temperature of the fluid (T_F) , the wire (T_W) and the heat input (Q) to the wire. Since the wire area was known, it was possible to calculate the heat transfer coefficient (h) as well as the fouling factor (R_F) and the fouling rate (dR_F/dt) .

Chemical analyses of materials used in this study including the Occidental desalted and high-pressure (second-stage) hydrotreated shale oil, the JP-8 derived from the hydrotreated oil, and the Arabian Light Berri petroleum desalted crude oil are shown in Table 35. Significant differences between the desalted petroleum and shale oils are the higher Bromine number, gravity and nitrogen content of the shale oil and the higher sulfur content in the petroleum oil. The hydrotreated Occidental shale oil, however, has a significantly lower Conradson carbon, Bromine number, iron, sulfur, nitrogen content and gravity than the original desalted oil.

Unfortunately, the state-of-the-art in predicting fouling has not advanced sufficiently to predict potential thermal fouling characteristics based on chemical analysis. However, as will be shown later, it is apparent that reduction in the sulfur and unsaturated compounds content as well as metals such as iron and nickel did result in lower fouling in the hydrotreated shale oil.

The thermal fouling characteristics of the Arabian Light Berri desalted petroleum crude and Occidental shale oil previously evaluated in June, 1980, were again reevaluated in March, 1981. Data from both tests are given in Tables 36 and 37, and in Figures 15 and 16.

The estimated age of the Occidental desalted shale oil by June, 1980 was 13 months, and 22 months by March, 1981. The Arabian Light Berri petroleum desalted crude was in storage about 8 months by June, 1980, and 17 months by March, 1981. Both oils were sparged with nitrogen when received and stored at 4.4°C (40°F) until ready for test. Prior to testing they were moved into a 40°C (105°F) room and stored there during the entire test period.

Discussion of Results

A regression analysis was made of the data and the fouling rate calculated for the 200° to 400° C (392° to 752° F) temperature range. As shown in Table 38, the aging of the Occidental desalted shale oil and the Arabian Light Berri petroleum crude oil significantly reduced fouling on the iron wire.

To illustrate potential effects of aging, the 250°C (482°F) fouling data, were plotted (Figure 17) assuming the fouling rate decreased linearly or exponentially. These data suggest that the dR_F/dt of the Occidental desalted slale oil had probably decreased by a factor of 5 to 10 times after aging at 4.4°C (40°F) for 22 months, and the Arabian Light Berri desalted petroleum crude decreased by a factor of 1.5 to 2 times after aging for 8 months. It is apparent, therefore, that more meaningful fouling data could be obtained with fresh samples.

Because it is anticipated that the shale oils will be blended with petroleum stocks for processing, the thermal fouling properties of blends of the Arabian Light Berri desalted crude containing 10 and 30% (by weight) of the Occidental desalted shale oil were determined. The data from these test are given in Tables 39 and 40 and presented graphically in Figures 18 and 19. When compared with the 100% petroleum and 100% shale oil results, it is apparent from examining Figure 20 that there were no significant differences in the thermal fouling characteristics of the blends. Blending did not result in the formation of any new fouling components, and the data actually show a minor decrease in fouling tendencies. These two oils, however, are quite old and the effect of blending should be repeated with fresh samples.

The thermal fouling properties of a high-pressure hydrotreated Occidental shale oil were determined and these data are shown in Table 41 and Figure 21. These data show that the high-pressure hydrotreated shale oil was a low fouling material with an activation energy significantly higher than that of the original desalted oil. Although specific compounds have not been identified, it can be speculated that this reduction in fouling is due to the reduction in concentration of unsaturated sulfur and nitrogen compounds. Fouling rate data calculated from a regression analysis of the data in Figure 21 for the 200° to 400°C (392°-752°F) region are given in Table 38.

The thermal fouling characteristics of a JP-8 derived from the high-pressure hydrotreated Occidental shale oil were determined and these data are summarized in Table 42 and Figure 22. These data show that the JP-8 had fouling properties greater than the hydrotreated shale oil but slightly lower than the original desalted oil. Apparently the catalytic process, converting the hydrotreated oil to the JP-8, results in the production of products which can foul thermally on a heated iron wire. No data were available to identify these products. Fouling rates calculated for the 200° to 400°C (392°-752°F) region are shown in Table 38.

In the Phase II report, the relative thermal fouling properties of two Paraho shale oils were reported. The fouling reactions did not apparently follow a typical Arrhenius concept, and it was speculated that these deviations were due to the presence of oxygenates. More recent experiments indicate that excessive fouling of the probe was responsible.

The test method requires that a build-up of fouling deposit has only a minor effect on the surface area or diameter of the original wire. It was shown that at temperatures from 250° to 311° C (482° - 592° F), a relatively long time of exposure of the iron wire at a constant voltage to Occidental desalted shale oil resulted in only minor changes in the fouling rate. The data from a similar set of experiments at 145° and 184° C(293° and 363° F) with the Paraho shale oil are summarized in Table 43 and Figure 23. It was obvious from the dramatic change in observed fouling rate that a build-up of deposit had occurred. Therefore, it was not possible with the high fouling materials to use the same experimental technique in which the probe temperature is increased during a run. A new probe must be used at each new temperature.

A new series of experiments were made with the Paraho shale oil employing a new probe for each new temperature (3 hours per test). Based on the data in Table 44 and shown in Figure 24, the new data are more reasonable and consistent than those obtained previously with the incremental increase in temperature technique. A regression analysis of the data was made, and the fouling rate in the 200° to 400°C (392°-752°F) range given in Table 38, confirms that the Paraho shale oil has a high fouling tendency.

During Phase II, Paraho shale oil was treated with a proprietary antifoulant resulting in a significant decrease in fouling rate. A second Paraho shale oil was also treated with the same proprietary antifoulant (40 ppm) and these fouling data, are given in Table 45 and Figure 24, also show a significantly reduced fouling rate.

Conclusions

The Monirex Fouling Monitor was used to determine the relative thermal fouling properties of aged Arabian Light Berri desalted petroleum, Occidental desalted shale oil and blends of these oils. In addition, the fouling properties of a hydrotreated Occidental shale oil, a JP-8 derived from this oil and a relatively new Paraho shale oil were also determined. These data, summarized in Table 38 and Figures 25 and 26, show that desalting and hydrotreating decrease the fouling tendency of shale oils, shale oils may have different fouling tendencies, and the aged Occidental shale oil had a fouling tendency similar to an Arabian Light Berri crude.

It was found that aging at 4.4°C (40°F), with relatively short periods at (40°C) 105°F, resulted in significant decreases in fouling rates. Since data are available for only two aging periods, it is difficult to project fouling rates to zero age. Although data are not available to show the fouling properties of a fresh sample, it can be speculated that an exponential decrease in fouling rate with aging indicates a rate 5 to 10 times higher for the fresh Occidental desalted shale oil than that found after 13 months aging, and up to 2 times greater for the Arabian Light Berri desalted petroleum oil when compared to those found after eight months. It is apparent, however, that the most meaningful data must be obtained with fresh samples.

Blending of the desalted Arabian Light Berri petroleum crude oil and the desalted Occidental shale oil at 70:30 or 90:10 ratios, by weight, had no significant effect on the fouling rates of either oil. These data indicate that blending did not result in the formation of any significant quantities of pro- or antifoulant materials. However, the activation energy of the fouling reactions of the blends were slightly higher than those of the original materials suggesting the possibility of a different fouling reaction mechanism. The original samples were quite old, with relatively low fouling rates, and it is apparent that these studies should be repeated with fresh oils.

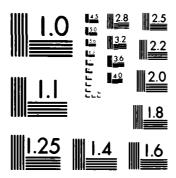
The high-pressure hydrotreated Occidental shale oil had a very low fouling rate. Although the analytical data show substantial removal of unsaturated sulfur, iron and nitrogen compounds by hydrotreating, specific compounds responsible for fouling have not been identified. The

activation energy of the hydrotreated oil fouling reaction is very high suggesting a different mechanism for the fouling reactions on the iron wire.

A JP-8 derived from the high-pressure hydrotreated Occidental desalted shale oil has a relatively low thermal fouling rate. This rate was slightly higher than that of the hydrotreated shale oil but lower than that of the original shale oil. Hydrocracking, the process used to make the JF-8, must apparently produce new precursors of fouling reactions as indicated by the reduction of the activation energy from 18 kcal to 6.5 kcal for the JP-8.

It was also shown that only low fouling materials such as the petroleum and the aged Occidental shale oil used can be evaluated with a single probe for all of the temperatures selected for fouling studies. A high fouling material, such as the relatively fresh Paraho shale oil, must be evaluated with a new probe at each new test temperature. Significant foulant build-up on the iron wire probe can change the wire surface area. Under these new conditions, it was shown that the Paraho shale oil fouling rate was approximately an order of magnitude higher than for the petroleum or Occidental shale oil.

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TABLE 35. ANALYSIS OF OCCIDENTAL DESALTED AND HYDROTREATED SHALE OIL,

JP-8 FROM HYDROTREATED OIL, PARAHO SHALE OIL

AND ARABIAN LIGHT BERRI DESALTED PETROLEUM OIL

	Light Berri Oil		Deashed 011	HP Hydrotreate 011	d JP-8	Paraho Shale Oil
API Gr. at 60°F	35.4		22.9	32.3	45.1	19.8
Sp. Gr. at 60°F	0.8478		0.9165	0.8639	0.8012	0.9352
Bubble Pt. at 500 psig, F	720		1056			
Distillation (D-1160), °F	,					
IBP	127		376	430	295	408
5%	225		467	469	314	471
10%	274		510	495	330	525
30%	420		621	581	359	670
50%	589		712	663	438	789
70%	782		820	760	474	889
87%	1083		953			1005
90%				958	512	
% Over	90		87	97	99	90
% Bottoms	10		13	3	••	10
Con. Carbon, wt-%	_		1.36	0.06	_	2.71
Nitrogen, wt-%	840	ppm	1.5	680 p	pm 1.1	ppm 2.2
Bromine Number	1.2		23.6	1.3	<u> </u>	34.7
Ash, wt-%	0.001		0.014	-	_	0.007
C7 Insol., wt-%	0.55		0.34	0.06	-	0.47
Emission, ppm						
Fe	0.55		42	1.3	-	38
Mn	-		<0.2	-	-	-
Cr	-		0.21	***	-	-
N1	2.4		6.7	0.25	-	2.2
Мо	-		1.6	<0.1	-	0.12
Cu	<0.1		<0.1	<0.1	-	0.18
H_2O , wt-%	323	ppm	0.05	-	-	<0.01
Arsenic, ppm	-		27.5	-	-	19
Sulfur, wt-%	1.34		0.64	0.012	282	
Total O, wt-%		ppm	0.65	-	-	1.58
Carbon, wt-%	85.74		84.99	85.8	84.3	-
Hydrogen, wt-%	12.96		12.27	13.6	13.9	-
Diene Value	~		31.6	-	-	-

TABLE 36. EFFECT OF WIRE TEMPERATURE ON h AND $dR_{\overline{F}}/dt$ VALUES OF AGED OCCIDENTAL DESALTED SHALE OIL

Period	Temperat Fluid	wire Wire	$\frac{\text{Wire Temp.}}{1/\text{K} \times 10^{-4}}$	h Original	Final	$\frac{dR_{F}/dt}{x \ 10^{-5}}$
			Run No. 35			
7	92.9	175.8	22.3	223.4	222.1	1.40
8	93.4	187.6	21.7	223.5	223.9	nil
9	93.4	211.2	20.7	227.9	225.0	1.45
10	93.4	235.6	19.7	226.7	222.9	2.52
11	93.4	264.2	18.6	223.1	220.8	2.24
12	93.4	294.6	17.6	218.6	213.1	4.43
13	93.4	327.9	16.6	212.2	204.1	6.08
14	93.4	365.9	15.7	201.5	194.5	6.70
15	93.4	406.8	14.7	191.2	178.8	12.9
16	93.4	458.6	13.7	174.8	155.6	26.7
17	93.9	526.4	12.5	153.1	137.4	30.5
			Run No. 37			
7	93.2	185.0	21.8	205.7	204.6	0
8	93.2	197.3	21.2	206.2	204.7	0.36
9	93.2	224.5	20.1	208.5	207.0	0.71
10	93.4	250.2	19.1	208.4	206.2	1.05
11	93.4	280.7	18.0	206.8	204.2	2.10
12	93.4	313.6	17.0	203.2	198.4	4.25
13	93.5	350.1	16.0	197.1	189.8	7.36
14	93.5	392.3	15.0	186.6	177.6	10.2
15	93.5	440.4	14.0	174.6	164.4	12.7
16	93.5	496.0	13.0	160.6	142.3	32.8
			Run No. 38			
7	92.0	182.4	21.9	208.6	208.6	0
8	92.0	194.4	21.3	210.3	210.3	0
9	92.0	220.2	20.2	214.3	212.6	0
10	92.0	246.0	19.2	213.6	212.3	0.74
11	92.0	275.9	18.2	212.0	210.7	1.70
12	92.0	307.8	17.2	209.4	205.7	2.67
13	92.5	341.9	16.2	204.7	199.8	4.50
14	92.5	380.1	15.3	197.0	190.9	6.13
15	92.5	421.1	14.4	188.8	181.9	6.89
16	92.7	466.4	13.5	179.2	172.2	8.19

TABLE 37. EFFECT OF WIRE TEMPERATURE ON h AND $dR_{\mbox{\scriptsize F}}/dt$ VALUES OF AGED ARABIAN LIGHT BERRI PETROLEUM CRUDE OIL

<u>Period</u>	Temperat Fluid	wire <u>Vire</u>	Wire Temp. $1/K \times 10^{-4}$	h Original	Final	$\frac{dR_{F}/dt}{\times 10^{-5}}$
			Run No. 33			
7	90.7	179.5	22.1	206.2	207.3	nil
8	91.1	190.6	21.6	209.8	210.7	nil
9	91.6	215.7	20.5	214.7	214.7	ni1
10	91.6	240.1	19.5	214.9	214.7	nil
11	92.0	268.2	18°•5	214.5	214.5	ni1
12	92.0	298.0	17.5	212.0	206.6	4.16
13	92.1	333.7	16.5	203.4	194.8	7.10
14	91.6	375.7	15.4	190.4	182.2	6.96
15	91.8	421.5	14.4	177.1	168.1	11.26
16	91.8	473.0	13.4	163.5	155.8	11.26
			Run No. 36			
7	90.8	182.1	22.0	196.4	197.5	nil
8	90.8	193.2	21.4	201.6	199.9	nil
9	90.9	218.3	22.2	204.6	205.0	ni1
10	91.1	242.8	19.4	205.8	205.8	nil
11	91.2	271.4	18.4	207.1	207.1	nil
12	91.4	300.0	17.4	206.9	205.5	1.18
13	91.7	330.9	16.6	204.8	200.5	3.81
14	91.7	365.6	15.7	198.9	193.8	5.00
15	91.8	403.6	14.8	191.3	181.7	8.24
16	91.8	449.0	13.8	178.9	171.8	9.05

TABLE 38. FOULING CHARACTERISTICS OF SHALE AND PETROLEUM PRODUCTS

					Fouling	Rate dR	<u>-/dt × 10</u>	5		
Temper	rature,	Occident	al Shale	Light		ALB-90	ALB-70		Paraho	JP-8 from
•c	<u>•</u> F	13 Mo	22 Mo	8 Mo	17 Mo	<u>05-10</u>	<u> 0S-30</u>	<u>HPOS</u>	3 Mo	<u>HPHOS</u>
200	392	3.49	1.07*	1.32	1.55*	0.59*	0.41*	0.009*	28	0.5
225	437	4.75	1.51	2.27	2.04*	0.90*	0.68*	0.02*	35	0.7
250	482	5.94	2.08	3.26	2.62*	1.32*	1.08*	0.06*	43	1.1
275	527	7.27	2.77	4.80	3.28*	1.87*	1.65*	0.13*	52	1.58
300	572	8.75	3.61	6.67	4.03	2.58	2.42	0.27*	62	2.23
325	617	10.37	4.61	9.01	4.87	3.46	3.45	0.51*	72	3,01
350	662	12.12	5.76	11.86	5.79	4.53	4.76	0.94*	83	4.07
375	707	14.00	7.07	15.30	6.80	5.81	6.43	1.65	95	5.27
400	752	15.97	8.55	19.34	7.87	7.30	8.46	2.77	107	6.53
Activa Ener	rgy									
cal.	/mol	-4626	-6613	-8176	-5142	-7797	-9605	-18011	-4214	-8550

^{* =} Calculated Values - Actual Values $dR_F/dt < 0.5 \times 10^{-5}$

ALB = Arabian Light Berri Petroleum Crude

OS = Occidental Shale Oil

HPHOS = High Pressure Hydrotreated Occidental Shale Oil

³ Mo. = Estimated Age of 3 Months

TABLE 39. EFFECT OF TEMPERATURE ON h AND dR_F/dt VALUES OF A 90:10 (BY WEIGHT) BLEND OF ARABIAN LIGHT BERRI DESALTED PETROLEUM CRUDE AND OCCIDENTAL SHALE OILS

Dowlad	Temperature, °C		$\frac{\text{Wire Temp.}}{1/\text{K} \times 10^{-4}}$	h		dR _F /dt x 10 ⁻⁵
Period	Fluid	Wire	1/K X 10	<u>Original</u>	Final	X 10 3
			Run No. 41			
7	91.3	178.7	22.1	209.2	208.7	nil
8	91.3	190.6	21.6	211.7	211.7	nil
9	91.6	215.6	20.5	214.5	214.2	nil
10	91.6	240.1	19.5	216.2	213.8	0.88
11	91.8	268.2	18.5	214.9	212.9	0.68
12	91.8	298.4	17.5	212.3	209.1	2.27
13	91.8	330.8	16.6	206.6	202.8	3.85
14	91.8	366.7	15.6	199.8	193.5	6.06
15	92.0	405.5	14.7	191.6	183.7	8.29
16	92.0	449.5	13.8	181.2	173.1	9.16
			Run No. 42			
7	90.7	178.0	22.2	213.6	213.6	nil
8	90.7	189.8	21.6	215.2	214.4	nil
9	90.9	215.0	20.5	217.3	216.6	ni1
10	90.9	239.9	19.5	216.5	215.9	1.23
11	90.9	269.2	18.4	214.3	214.2	nil
12	91.1	299.1	17.5	212.6	211.3	1.78
13	91.3	331.8	16.5	207.6	202.4	3.52
14	91.6	367.9	15.6	200.7	188.1	10.71
15	91.6	415.5	14.5	184.7	174.1	10.51
16	91.6	464.9	13.6	171.1	166.0	8.06

TABLE 40. EFFECT OF TEMPERATURE ON h AND dR_F/dt VALUES OF A 70:30 (BY WEIGHT) BLEND OF ARABIAN LIGHT BERRI DESALTED PETROLEUM AND OCCIDENTAL DESALTED SHALE OILS

Period	Temperat Fluid	ure, °C Wire	Wire Temp. $1/K \times 10^{-4}$	h Original	<u>Final</u>	$\frac{dR_{F}/dt}{x \ 10^{-5}}$
			Run No. 39			
7	91.6	176.4	22.2	219.8	219.8	nil
8	91.8	187.9	21.7	220.8	220.8	nil
9	91.8	213.0	20.6	222.4	222.3	nil
10	91.8	237.1	19.6	222.2	222.0	nil
11	91.8	264.8	18.6	221.4	220.9	0.85
12	91.8	293.5	17.6	219.9	219.6	0.70
13	91.8	324.3	16.7	216.8	213.3	3.22
14	92.1	358.7	15.8	209.5	203.8	4.71
15	92.3	399.4	14.9	199.5	182.5	15.17
16	92.3	450.2	13.8	181.8	172.1	10.16
			Run No. 40			
7	91.3	178.4	22.2	211.9	213.8	nil
8	91.3	189.5	21.6	213.9	215.0	nil
9	91.8	214.1	20.5	218.7	218.2	nil
10	91.6	238.4	19.6	218.5	218.5	nil
11	91.6	266.0	18.6	218.1	218.1	nil
12	91.7	294.4	17.6	217.6	215.7	1.56
13	91.7	326.1	16.7	213.1	208.1	3.85
14	91.7	364.2	15.7	202.9	194.7	7.89
15	91.8	406.8	14.7	190.6	178.6	12.16
16	91.8	456.1	13.7	176.0	168.5	9.76

TABLE 41. EFFECT OF PROBE TEMPERATURE ON h AND $dR_{\mbox{\scriptsize F}}/dt$ OF HIGH PRESSURE HYDROTREATED OCCIDENTAL SHALE OIL

Period	Temperat Fluid	wire	Wire Temp. 1/K x 10 ⁻⁴	h Original	Final	$\frac{dR_{F}/dt}{\times 10^{-5}}$
			Run No. 47			
7	91.1	174.0	22.4	226.8	226.8	nil
8	91.6	184.8	21.8	231.0	231.0	nil
9	91.6	209.4	20.7	231.7	231.7	nil
10	91.6	232.9	19.8	231.9	231.9	nil
11	92.0	261.0	18.7	230.0	230.0	nil
12	92.0	288.5	17.8	229.0	229.0	nil
13	92.0	318.1	16.9	226.6	226.6	nil
14	92.5	348.8	16.1	223.8	223.8	nil
15	92.5	380.4	15.3	220.4	218.3	1.94
16	92.5	414.9	14.5	214.7	208.6	4.23
			Run No. 49			
7	90.7	183.1	21.9	193.4	193.4	nil
8	90.7	194.3	21.4	197.0	197.0	nil
9	90.7	219.6	20.3	201.0	201.0	nil
10	91.1	243.9	19.3	203.9	203.9	nil
11	90.7	270.4	18.4	206.1	206.1	nil
12	90.7	297.8	17.5	208.1	208.1	nil
13	90.7	328.8	16.6	205.2	205.2	nil
14	91.1	359.4	15.8	204.2	204.2	nil
15	92.0	390.1	15.1	203.9	201.9	1.97
16	92.0	425.1	14.3	199.5	193.9	4.70
17	91.6	466.1	13.5	190.5	181.8	9.07
18	91.6	515.9	12.7	177.3	171.8	8.71

TABLE 42. EFFECT OF WIRE TEMPERATURE ON h AND dR_F/dt OF JP-8 DERIVED FROM HIGH-PRESSURE HYDROTREATED OCCIDENTAL SHALE OIL

Period	Tempera Fluid	ture, °C Wire	$\frac{\text{Wire Temp.}}{1/\text{K} \times 10^{-4}}$	h Original	<u>Final</u>	$\frac{dR_{F}/dt}{x 10^{-5}}$
			Run No. 64			
5	96.1	155.9	23.3	169.1	169.1	nil
6	96.2	178.2	22.2	179.1	180.4	nil
7	96.3	193.0	21.5	183.4	182.0	0.26
8	96.3	205.9	20.9	184.1	186.6	nil
9	96.4	232.7	19.8	188.2	188.4	nil
10	97.0	258.4	18.8	189.7	189.7	nil
11	97.5	287.4	17.8	191.7	190.0	1.62
12	98.2	319.7	16.9	189.8	186.1	3.20
13	98.4	353.9	16.0	155.5	152.1	4.35
14	98.6	389.8	15.1	181.7	177.6	4.15
			Run No. 65			
7	100.2	251.2	19.1	211.1	211.1	nil
8	100.5	280.5	18.1	210.2	208.4	1.81
9	100.5	311.7	17.1	207.6	205.0	2.46
10	100.5	344.5	16.2	203.1	198.1	3.29
11	100.5	380.0	15.3	197.3	193.2	3.98
12	100.8	417.3	14.5	191.8	186.9	4.67
13	100.8	456.7	13.7	184.9	174.0	11.74
14	100.8	504.2	12.9	174.2	160.2	16.75

TABLE 43. EFFECT OF TIME AT CONSTANT WIRE VOLTAGE ON h AND $dR_{\mbox{\scriptsize F}}/dt$ OF PARAHO SHALE OIL

Period, hours	Tempera Fluid	ture, °C Wire	$\frac{\text{Wire Temp.}}{1/\text{K} \times 10^{-4}}$	h Original	Final	dR _F /dt x 10 ⁻⁵
			Run No. 60			
0-3	92.4	183.8	21.9	234.8	189.2	33.6
3-6	92.4	201.1	21.1	188.2	168.5	20.3
6-9	92.4	211.6	20.6	167.1	153.5	18.0
9-12	92.3	219.8	20.3	153.1	141.4	17.7
12-15	92.4	227.8	20.0	140.7	130.8	16.6
15-18	89-95	240.0	19.5	127.5	119.8	19.7
18-21	89-95	245.3	19.3	121.6	116.7	17.0
21-24	92.3	248.6	19.2	116.1	112.1	14.0
0-24	92.4	183.8	21.9	234.8	112.1	19.4
			Run No. 61			
0-3	92.6	144.9	23.9	199.0	167.6	26 E
3-6	92.6	153.2	23.5	167.6	154.2	26.5
6-9	92.7	157.2	23.2	154.2	148.2	15.1 10.4
				22.02	14002	10.4
9-12	92.6	159.4	23.1	148.2	141.8	8.23
12-15	92.6	161.8	23.0	142.3	139.4	5.58
15-18	92.7	162.9	22.9	139.4	136.1	2.15
18-21	92.8	164.3	22.9	136.1	134.5	2.77
21-24	92.8	165.7	22.8	132.9	131.4	6.48
24-27	92.8	166.9	22.7	131.3	130.2	4.09
27-30	92.8	167.3	22.7	130.2	128.4	2.13
30-33	92.7	168.3	22.7	128.2	127.1	3.22
33-36	92.8	168.7	22.6	127.1	126.5	0.70
36-39	91.6-92.9	168.7	22.6	126.5	125.4	nil
39-42	92.7	169.8	22.6	124.6	124.2	nil
42-45	92.8	170.2	22.6	124.2	124.2	nil
45-48	92.8	170.2	22.6	124.2	122.6	3.65
48-51	92.9	170.6	22.5	122.6	122.8	nil
51-54	92.8	170.9	22.5	122.8	121.7	2.94
54-57	92.7	171.3	22.5	121.7	120.7	1.89
57-60	92.8	171.7	22.5	120.7	120.7	1.64
0-60	92.7	144.9	23.9	199.0	120.7	5.43

TABLE 44. EFFECT OF TEMPERATURE ON h AND $dR_{\rm F}/dt$ VALUES OF PARAHO SHALE OIL. NEW WIRE PROBE AT EACH NEW TEMPERATURE

	Temperat	ture, °C	Wire Temp.	ħ		dR _F /dt
Period	Fluid	Wire	$1/K \times 10^{-4}$	Original	Final	$\times 10^{-5}$
58	92.0	143.0	24.0	203.4	191.5	11.9
61	92.5	145.8	23.9	199.0	167.6	26.5
62	91.1	148.8	23.7	179.8	164.2	17.4
56	92.5	165.2	22.9	206.8	188.0	15.8
60	92.4	183.8	21.9	234.8	189.2	33.6
55	91.8	186.8	21.8	226.9	202.6	16.8
54	91.8	188.7	21.7	220.6	194.5	18.9
53	91.6	190.9	21.5	206.6	170.1	32.8
57	97.2	213.0	20.6	224.8	194.8	25.1
59	91.6	235.8	19.6	223.2	162.5	52.6

TABLE 45. EFFECT OF PROPRIETARY ANTIFOULANT ON FOULING RATE AND h VALUES OF PARAHO SHALE OIL

dR _F /dt		h	Wire Temp.	ure, °C	Temperat	
$\times 10^{-5}$	Final	Original	$1/K \times 10^{-4}$	Wire	Fluid	Period
nil	237.1	235.4	22.4	173.0	92.2	10
ni1	241.9	236.5	21.9	183.3	92.6	11
nil	245.8	241.2	20.9	206.2	92.3	12
11.97	228.8	244.6	20.0	228.2	92.3	13
17.5	203.7	225.6	18.6	263.6	92.5	14

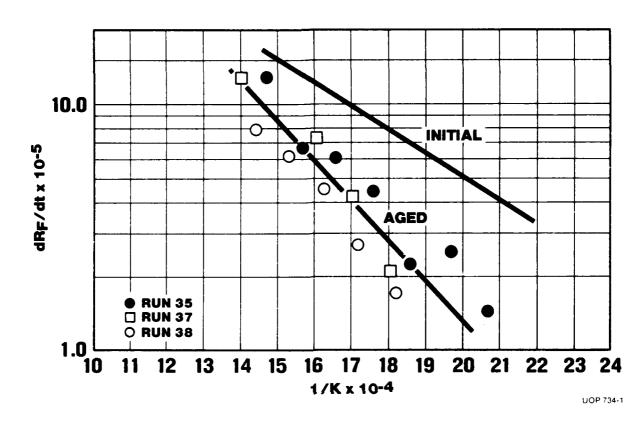
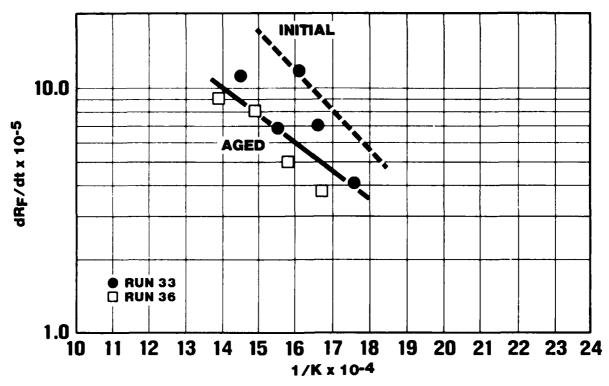


FIGURE 15
EFFECT OF TEMPERATURE AND AGING ON FOULING RATE OF DESALTED OCCIDENTAL SHALE OIL



UOP 734-2

FIGURE 16
EFFECT OF TEMPERATURE AND AGING ON
FOULING RATE OF DESALTED ARABIAN LIGHT
BERRI PETROLEUM OIL

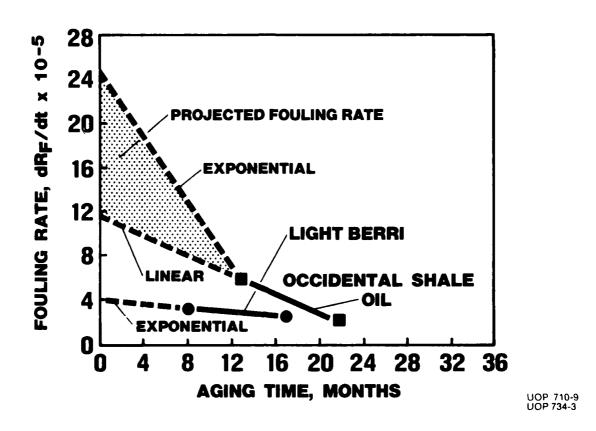
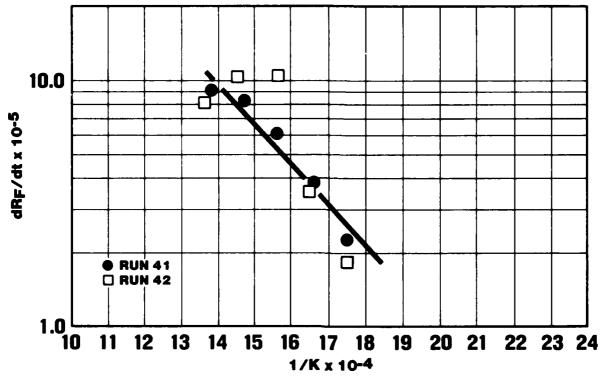


FIGURE 17
EFFECT OF AGING ON FOULING
RATE



UOP 734-4

FIGURE 18

FOULING RATE OF 1:9 BLEND OF DESALTED OCCIDENTAL SHALE AND ARABIAN LIGHT BERRI PETROLEUM OILS

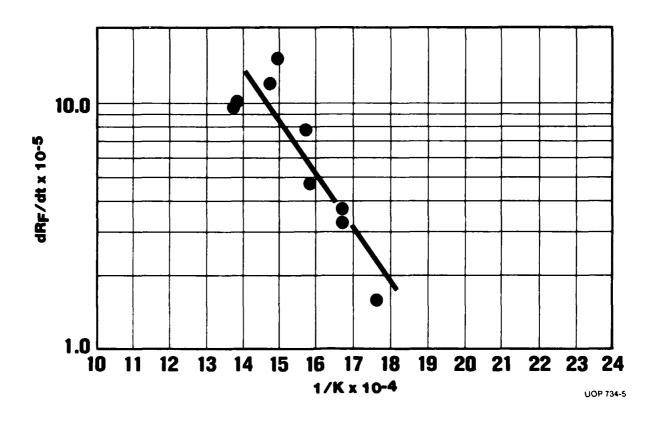


FIGURE 19
FOULING RATE OF 3:7 BLEND OF DESALTED OCCIDENTAL SHALE AND ARABIAN LIGHT BERRI PETROLEUM OILS

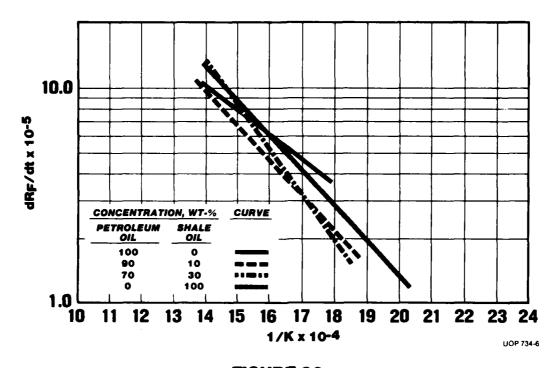


FIGURE 20
EFFECT OF BLENDING ON FOULING RATE

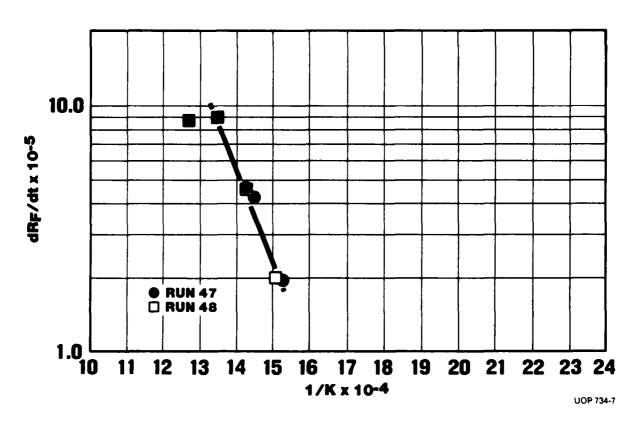


FIGURE 21
FOULING RATE OF HYDROTREATED
OCCIDENTAL SHALE OIL

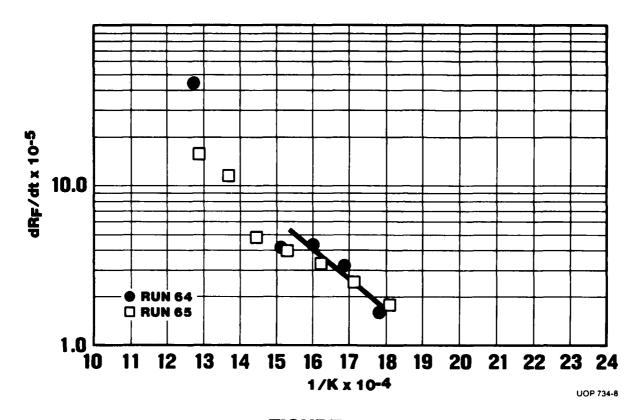


FIGURE 22
FOULING RATE OF JP-8 FUEL
FROM HYDROTREATED SHALE OIL

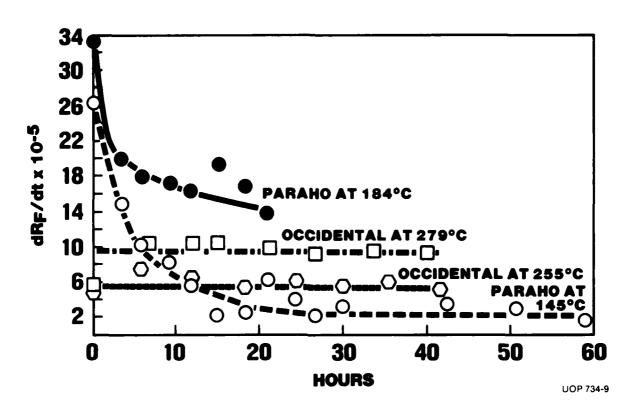


FIGURE 23
FOULING RATES OF DESALTED
OCCIDENTAL AND PARAHO
SHALE OILS

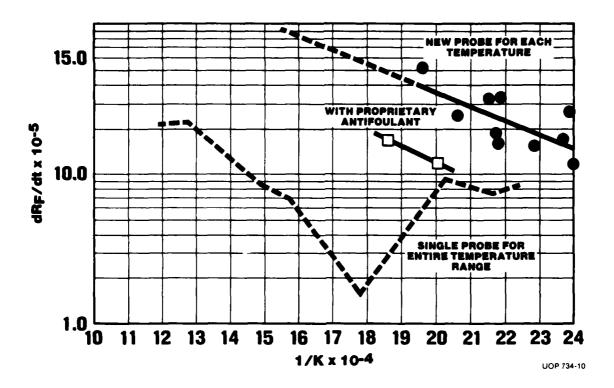


FIGURE 24
FOULING RATE OF PARAHO SHALE OIL

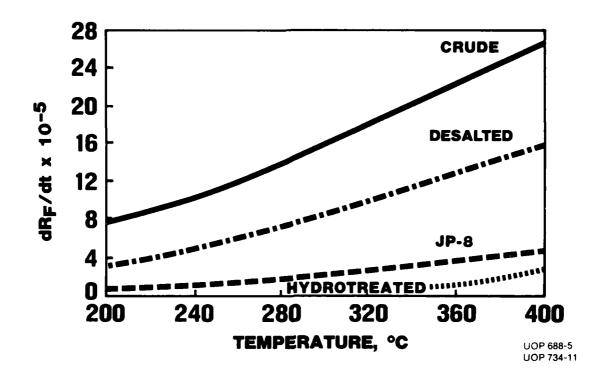


FIGURE 25

EFFECT OF TEMPERATURE ON THE FOULING RATE OF OCCIDENTAL SHALE OIL DERIVED FEEDS

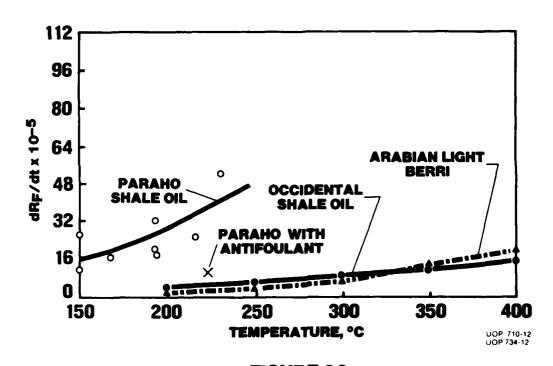


FIGURE 26
EFFECT OF TEMPERATURE ON FOULING
RATE OF SHALE AND PETROLEUM OILS

SECTION VII

SHALE OIL/PETROLEUM OIL STABILITY/COMPATIBILITY STUDY

Introduction

Background

Oil shale represents an enormous potential reserve of energy for the United States. Conversion of oil shale to liquid hydrocarbons suitable as substitutes for transportation and heating fuels could augment the United States' reserves. Various pyrolysis processes are presently under development to produce liquid hydrocarbons from oil shale. They include surface processes and "in-situ" processes. Several processes have progressed beyond the small bench scale of the laboratory to the demonstration stage, and now have the capability of producing relatively large quantities of hydrocarbon liquids. Characterization of the liquid generated from these processes is an important consideration in direct utilization or upgrading processes. The immediate and best utilization of hydrocarbon liquids produced from demonstration plants and commercial prototypes may be blending them with petroleum crude oils or fractions thereof and processing the blends in existing refineries.

Objectives

The objectives of this study were to determine the stability of primary shale oils and treated products and the compatibility/stability of primary shale oils and treated products with petroleum crude oil. The stability of shale oils and treated products in storage and use is of concern to refiners and users. Whereas many studies have been made on the stability of petroleum liquids, stability of shale oils and treated products have been limited by unavailability of shale oils. Their resistances to chemical change and physical disintegration in storage is not well known. The compatibility of shale oils and treated products with petroleum rude oil is also of concern. The compatibility of shale oils and treated products with petroleum rude oil is an important consideration in

their blending, particularly with respect to co-processing shale oil/petroleum crude oil in existing refineries. The resulting blends should form a homogeneous mixture that neither separates nor is altered by chemical interaction. The physical and chemical properties of the blend should not be adversely affected.

Outline of Program

The program involved the stability study of primary shale oil and treated products derived therefrom and the compatibility/stability of primary shale oils and treated products derived therefrom with petroleum crude oil.

The primary shale oils selected were a raw Paraho shale oil and a raw Occidental shale oil. The crude oil selected was a heavy Arabian crude. Treated shale oils were derived from single-stage and two-stage hydrotreatment of the primary shale oils. Compatibility/stability studies of the shale oil/crude oil blend were conducted on blends consisting of 30 volume percent shale oil. The primary shale oils, treated products and petroleum crude oil were deoxygenated and inspected before the blending and subsequent stability testing.

Compatilibity and Stability Studies

Sources and Properties of Primary Samples

Primary Shale Oils -- Paraho shale oil produced by direct fired retort and Occidental shale oil produced by in-situ retorting were selected for study. The Paraho shale oil assigned for work under Phase II of the contract was received in March, 1979. It was from the 100,000 barrels of crude Paraho shale oil produced by Paraho Development Corporation over the three-year period from 1976 to 1978. The raw Paraho shale oil was desalted and dewatered. Inspections are given in Table 46. The Occidental shale oil, also processed under the aforementioned contract, was received in April, 1979. It was from the underground Room 6 retort using Occidental Petroleum Company's Modified In-Situ retorting process.

The raw Occidental shale oil was desalted and dewatered. Inspections are given in Table 46.

<u>Crude Oil</u> -- The foreign crude selected to represent typical Gulf Coast processing status was a heavy Arabian crude. Inspections are given in Table 46.

Treatment and Properties of Hydrotreated Shale Oils

Shale Oils -- Shale oils have unique characteristics relative to conventional crude petroleum oil. Unusually high arsenic and iron levels, high pour point and viscosity, a high unsaturates concentration, and a significant solids (ash) content make conventional front-end refining techniques unusable without proper pretreatment. To prepare the shale oil for the primary conversion step -- hydrocracking to jet and other fuels -- two stages of hydrotreatment were selected. The stability of the hydrotreated products and the stability/compatibility of petroleum crude oils blends were studied.

<u>First-Stage Hydrotreatment</u> -- The desalted Paraho shale oil and the desalted Occidental shale oil were processed in a first-strage hydrotreater to provide for removal of certain contaminants and stabilization of the raw shale oil, to allow high severity hydrotreatment and subsequent conversion operations. Inspections are given in Table 46.

Second-Stage Hydrotreatment -- The Paraho and Occidental first-stage hydrotreated products were processed in a second-stage hydrotreater at higher severity to reduce the high levels of impurities such as nitrogen and oxygen to acceptable levels for the primary conversion step -- hydrocracking to jet and other fuels. Inspections are given in Table 46.

Experimental Procedures

The preparation and analysis of samples for stability and compatibility testing are described in Appendicies A.1 and A.2. The stability tests selected for study were the Three Months 110°F Dark Storage Fuel Oil Stability Test (See Appendix A.3), the E. I. DuPont de Nemours and Co.

300°F Accelerated Fuel Oil Stability Test (1) and the ASTM No. 2274-74 Oxidation Stability of Distillate Fuel Oil Test (2). These tests are applicable to petroleum distillate fuels and were modified to permit stability testing of shale oils, petroleum crude oils and blends thereof (See appendices A.4, A.5 and A.6).

Results and Discussion

Stability Studies

Stability was assessed by measuring formation of adherent insolubles and changes in heptane insolubles, toluene insolubles and viscosity. The results obtained are summarized in Tables 47 and 48 and Figures 27 through 30.

Adherent Insoluble Formation — The samples show relatively good stability with respect to adherent insoluble formation with the possible exception of the 30 volume percent raw Paraho shale oil/heavy Arabian crude blend which had adherent gum values of 2.1 mg/100 mL in the three-months storage test and 10.6 mg/100 mL in the ASTM D 2274-74 test. The other adherent gum values were less than 1 mg/100 mL in the three-month storage test.

<u>C7 Insoluble Stability</u> -- The second-stage hydrotreated Paraho shale oil and the first- and second-stage Occidental shale oils were stable with respect to C7 insolubles.

⁽¹⁾ Petroleum Laboratory Mechod No. F21-61, "300°F Accelerated Fuel Oil Stability Test", E.I. Dupont de Nemours and Company, Inc., Wilmington, Delaware, December, 1962.

⁽²⁾ Annnual Book of Standards, Part 24, P. 296, 1977; American Society for Testing and Materials, 1916 Race Street, Philadelphia, Pennsylvania 19103.

The C7 insolubles of the heavy Arabian crude, the raw and first-stage hydrotreated Paraho shale oil and raw Occidental shale oil increased after stability testing, indicating some measure of instability with respect to C7 insolubles. The heavy Arabian crude had the highest increase indicating it was the most unstable. Next were the raw Paraho and raw Occidental shale oils which had similar increases. The first-stage hydrotreated Paraho had the lowest increase. Stability with respect to C7 insolubles improved with the degree of hydrotreatment.

Blending of the raw shale oil with heavy Arabian crude decreases stability with respect to C7 insolubles as evidenced by the magnitude of the increase in the C7 insolubles after the three-month storage and the ASTM D 2274-74 test. Blending of the first- and second-stage hydrotreated shale oil with the heavy Arabian crude improves stability with respect to C7 insolubles as evidenced by the decrease in the magnitude of the increase. Stability of the blends improves with the degree of hydrotreatment.

Viscosity Stability -- The second-stage hydrotreated Paraho shale oil and Occidental shale oil were stable with respect to viscosity. viscosity of the heavy Arabian crude, raw and first-stage hydrotreated Paraho shale oil, and raw and first-stage hydrotreated Occidental shale oil increased after stability testing, indicating some measure of instabil-The heavy Arabian crude had the highest increase indicating it was the most unstable. Next were the raw Paraho and Occidental shale oils which had similar increases. The lowest increases were obtained by the first-stage hydrotreated Paraho and first-stage hydrotreated Occidental shale oil which had similar increases. Stability with respect to viscosity improves with the degree of hydrotreatment. Blending of the heavy Arab'an crude oil with the shale oils improves stability with respect to Stability of the blends improves with viscosity. the hydrotreatment.

<u>Toluene Insoluble Stability</u> -- All samples show good stability with respect to toluene insolubles.

Compatibility Studies

The compatibility of the blends was determined by comparing the experimental values of viscosity, C7 insolubles and toluene insolubles of the blend to calculated values. The calculated values of viscosity were determined from viscosity blending charts (3, 4). The calculated values of C7 insolubles and toluene insolubles were determined from the amount of insolubles present in the original samples. The results obtained are summarized in Table 49. Good agreement was obtained between the calculated and experimental values in all the tests indicating that all of the blends of 30 volume percent shale oil/heavy Arabian crude oil are compatible.

Conclusions

The raw shale oils and hydrotreated products were found to be compatible with the heavy Arabian crude oil.

The raw shale oils, hydrotreated products and heavy Arabian crude showed good stability with respect to toluene insolubles and adherent insolubles.

The second-stage hydrotreated shale oils were found to be stable with respect to C7 insolubles and viscosity.

The first-stage hydrotreated Occidental shale oil was found to be stable with respect to C7 insolubles.

The heavy Arabian crude showed some measure of instability with respect to C7 insolubles and viscosity and was the most unstable of the materials tested.

⁽³⁾ UOP Calculation Charts -- RE Chart I-4.

^{(4) &}quot;Refutas" Viscosity-Temperature Chart Designed by C. I. Kelly, Copyright BAIRO and TATLOCK (London), Ltd.

The raw shale oils showed some measure of instability with respect to C7 insolubles and viscosity but less than the heavy Arabian crude.

The first-stage hydrotreated Paraho shale oil showed some measure of instability with respect to C7 insolubles and viscosity but less than the raw shale oil.

The first-stage hydrotreated Occidental shale oil showed some measure of instability with respect to viscosity but less than the raw shale oil.

Stability with respect to C7 insolubles and viscosity improves with increased hydrotreatment.

Blending of the shale oils with heavy Arabian crude improves stability with respect to viscosity.

Blending of the raw shale oils with heavy Arabian crude decreases stability with respect to C7 insolubles.

Blending of the first- and second-stage hydrotreated shale oil with the heavy Arabian crude improves stability with respect to C7 insolubles. Stability of blends with respect to C7 insolubles and viscosity improves with increased hydrotreatment. Instability reflected in the shale oil/petroleum crude oil blends is accounted for to a great extent by the instability of the petroleum crude oil and should have no adverse effects in co-processing.

The data obtained were derived from shale oils that were stored for two to four years before the compatiblity and stability tests were conducted and may not be representative of data derived from fresh commercial samples.

TABLE 46. INSPECTION OF OILS

	Heavy		· ·	:			
	Arabian		Parano shale UI	_		Occidental Shale Ull	9 011
	Crude	Raw	First-Stage	Second-Stage	Raw	First-Stage	Second-Stage
			Hydrotreated	Hydrotreated		Hydrotreated	Hydrotreated
API Gravity at 60°F	26.4	25.2	30.3	33.5	22.5	26.2	32.2
Specific Gravity at 60°F	0,8961	0,9303	0.9111	0.8576	0.9188	0.8973	0.8644
Distillation (D-1160), °F							
18P	138	372	390	344	399	422	356
55	189	450	471	438	479	508	450
10%	290	200	517	471	512	538	488
20%	420	585	290	521	572	589	532
30%	518	999	651	574	623	631	570
40%	626	728	710	620	119	672	615
50\$	735	784	160	670	729	718	099
809	851	840	810	721	775	191	710
70\$	176	888	861	165	822	812	160
80\$	(76) 1039	950	919	821	878	872	810
\$ 06		1025	(95) 1043	(EP) 1009	096	(94) 1010	6FP) 979
S Over	76.0	0.06	95.0	0.66	0.06	94.0	97.0
Bottoms	24	10.0	5.0	0.1	10.0	0.9	3.0
Carbon, wt-%	80.88	84.6	85,28	85.93	84.59	85.02	85,73
Hydrogen, wt-≴	11.22	11.4	12,05	12,40	12.42	12.60	13.01
Oxygen, wt-%	0.93	1.76	09*0	0.26	1.55	0.52	0.28
Sulfur, wt-X	2,97	0.71	0.0746	0,00323	0.68	0.0261	< 0.03
Nitrogen, wt-%	0.17	2.19	1,90	90.0	1.42	1.09	60°0
Cy insolubles, wt-\$	4.67	0.44	0.14	0.01	0.26	0.01	0.01
Toluene Insolubles, wt-\$	< 0.01	0.01	< 0.01	0.01	< 0.01	0.01	< 0.01
Viscosity at 100°F, SUS	118.5	213	156.0	54.2	186.7	124.6	60.1
Viscosity at 100°F, cSt	24.80	45.72	33,23	8,633	39.98	26.20	10.35
Steam Jet Gum, mg/100 mL	35,875.0	26,463.0	22,129.0	5,668.1	21,314.8	16,940.5	7,823.5
Bromine Number	2.4	30.2	15.8	0 •8	19.4	7.8	1.9
Nickel, ppm	51	3,5	3.0	۰ 0 م	8.8	3.2	< 0.1
Vanadium, ppm	47	0.5	0.5	< 0.1	0.5	< 0.2	< 0.2
Iron, ppm	2.8	108	IJ	< 0.3	53	7	< 0.1
Arsenic, ppm	~	30	- •	~	28	- •	- ~
Conradson Carbon, wt-%	7,98	2,35	1,57	0.01	1.05	0.44	0.03

TABLE 47. PARAHO SHALE OIL-PETROLEUM OIL

Compatibility/Stability Study

Nате	Heavv	^	30% Paraho Shale 011/ 70% Heavy Arabian Crude	011/ Crude		Day Change	-
	Arabian		First-Stage	Second-Stage		First-Stage	Second-Stage
	Crude	Raw	Hydrotreated	Hydrotreated	Raw	Hydrotreated	Hydrotreated
Initial Analysis:							
C7 insolubles, wt-\$	4.67	3,23	3.24	3,30	0.44	0.14	0.01
Toluene insolubles, wt-8	< 0.01	< 0.01	0.01	< 0.01	0.01	< 0.01	0.01
Viscosity at 100°F, SUS	118.5	130.6	124.5	86.9	213	156	54.2
Stability Test Analysis:							
Du Pont F-21-61							
C7 insolubles, wt-%	5.92	3,38	3,28	3,60	0.40	0.2	< 0.01
Toluene insolubles, wt-\$	0.07	0.01	0.01	< 0.01	< 0.01	0.01	0,01
Viscosity at 100°F, SUS	190.9	178.9	162,1	111.5	268	183.7	56.3
D-2274 16-hr Heat Treatment							
Cy Insolubles, wt-\$	6.23	5.17	4.23	4.19	1.70	0,79	0,02
Toluene insolubles, wt-%	< 0.01	0.01	< 0.01	< 0.01	< 0.01	0.01	< 0,01
Viscosity at 100°F, SUS	214	246	194.3	115.9	325	205	55.4
Adherent Gum, mg/100 mL	1.2	10.6	2.2	6*0			•
3-months Dark Storage, 110°F							
C7 insolubles, wt-8	6.07	4.48	4.12	4.14	0.97	0,29	0.01
Toluene insolubles, wt-\$	0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.01	< 0.01
Viscosity at 100°F, SUS	381	358	286	115.4	311	171	55.5
Adherent Gum, mg/100 mL	0.5	2.1	0.3	0.2	0.5	0.8	0.8

TABLE 48. OCCIDENTAL SHALE OIL/PETROLEUM OIL

Compatibility/Stability Study

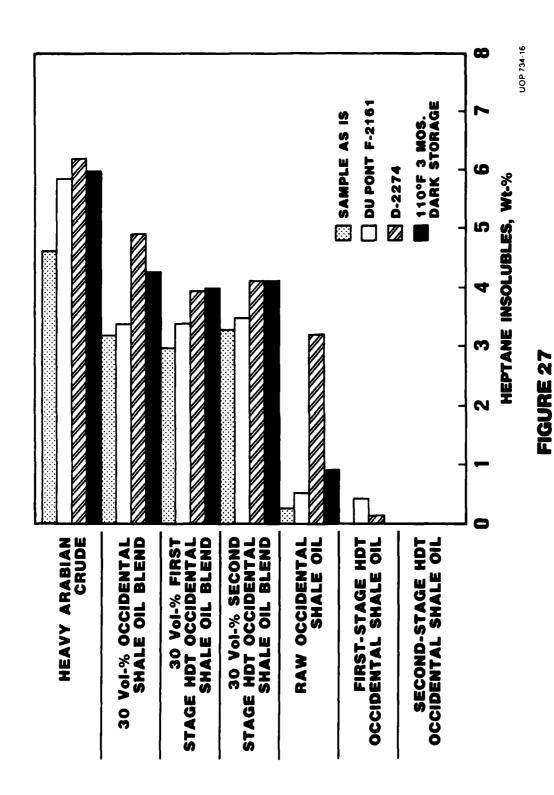
Мале	Неачу	*1	30% Occidental Shale Oil/ 70% Heavy Arablan Crude	516 011/ Crude		Occidental Shale Oll	le 011
	Arablan		First-Stage	Second-Stage		First-Stage	Second-Stage
	Crude	Raw	Hydrotreated	Hydrotreated	Raw	Hydrotreated	Hydrotreated
initial Analysis:							
Cy Insolubles, wt-\$	4.67	3,21	3,00	3,28	0.26	0.01	0.01
Toluene Insolubles, wt-5	< 0.01	< 0.01	0.02	0.01	< 0.01	0.01	< 0.01
Viscosity at 100°F, SUS	118.5	132.5	115.6	89.7	186.7	124.6	60.1
Stability Test Analysis:							
Du Pont F-21-61							
Cy Insolubles, wt-\$	5,92	3,38	3,39	3,51	0.50	0.40	0.01
Toluene Insolubles, wt-\$	0.07	< 0.01	0.01	< 0.01	0.01	0.01	0.01
Viscosity at 100°F, SUS	190.9	169.7	146.7	107.9	210	135	63.6
D-2274 16-hr Heat Treatment							
Cy insolubles, wt-%	6.23	4.95	3,94	4.17	3,18	0.11	< 0.01
Toluene Insolubles, wt-\$	< 0.01	< 0.01	< 0.01	< 0.01	0.01	< 0.01	< 0.01
Viscosity at 100°F, SUS	214	209	169.5	119.4	260	140.1	62,2
Adherent Gum, mg/100 mL	1.2	2.4	0.4	1.7		6*0	-:
3-months Dark Storage, 110°F							
Cy insolubles, wt-%	6.07	4.68	3.97	4.17	0.88	0.03	< 0.01
Toluene insolubles, wt-\$	0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Viscosity at 100°F, SUS	381	329	244	163.5	221	131.6	59.7
Adherent Gum, mg/100 mL	0.5	6.0	0.3	0.3	7.0	0.1	0.2

TABLE 49. SHALE OIL COMPATIBILITY/STABILITY STUDY

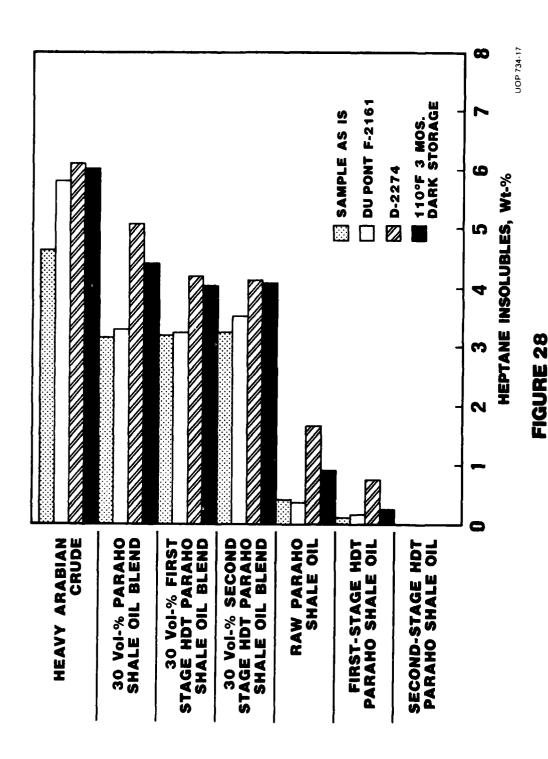
Comparison of Analyzed and Calculated Values for Shale Oil and Petroleum Blends

	Analytical	Calc	ulated
	cSt Viscosity	cSt Vi	scosity
Shale Oil Component in Blend		RE 1-4	Refutas
30 Vol-% Raw Paraho Shale 011	27.54	29.2	29.66
30 Vol-% Paraho Shale Oil First-Stage Hydrotreated	25.17	26.9	27.03
30 Vol-% Paraho Shale Oil Second-Stage Hydrotreated	17.38	17.5	17.59
30 Vol-% Raw Occidental Shale Oil	27.97	28.0	28.55
30 Vol-% Occidental Shale Oil First-Stage Hydrotreated	24.15	25.2	25.2
30 Vol-% Occidental Shale Oil Second-Stage Hydrotreated.	18.06	18.8	18.73

	Ana	lytical	Calcu	lated
	C ₇ Insol., Wt-%	Toluene Insol., Wt-%	C7 Insol., Wt-%	Toluene Insol., Wt-%
30 Vol-% Raw Paraho Shale Oil	3.23	<0.01	3.36	<0.01
30 Vol-% Paraho Shale Oil First-Stage Hydrotreated	3.24	0.01	3.29	<0.01
30 Vol-% Paraho Shale Oil Second-Stage Hydrotreated	3.30	<0.01	3.31	<0.01
30 Vol-% Raw Occidental Shale Oil	3.21	<0.01	3.32	<0.01
30 Vol-% Occidental Shale Oil First-Stage Hydrotreated	3.00	0.02	3.27	<0.01
30 Vol-% Occidental Shale Oil Second-Stage Hydrotreated	3.28	<0.01	3.31	<0.01

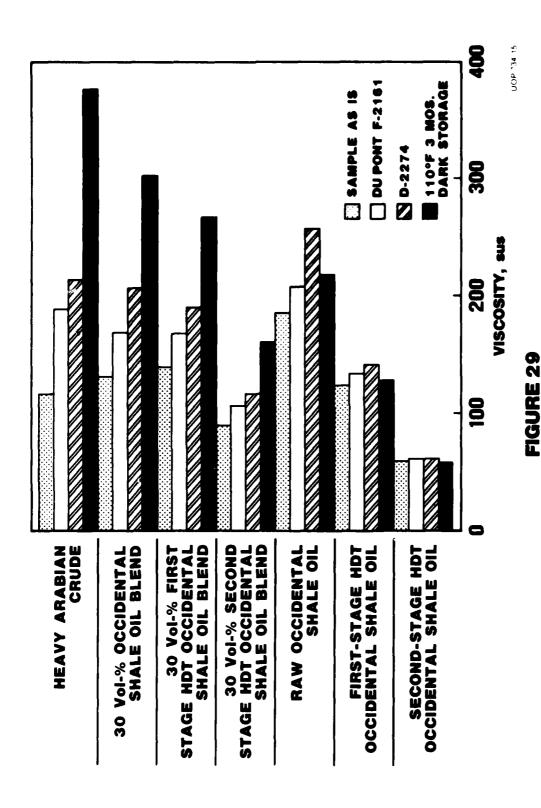


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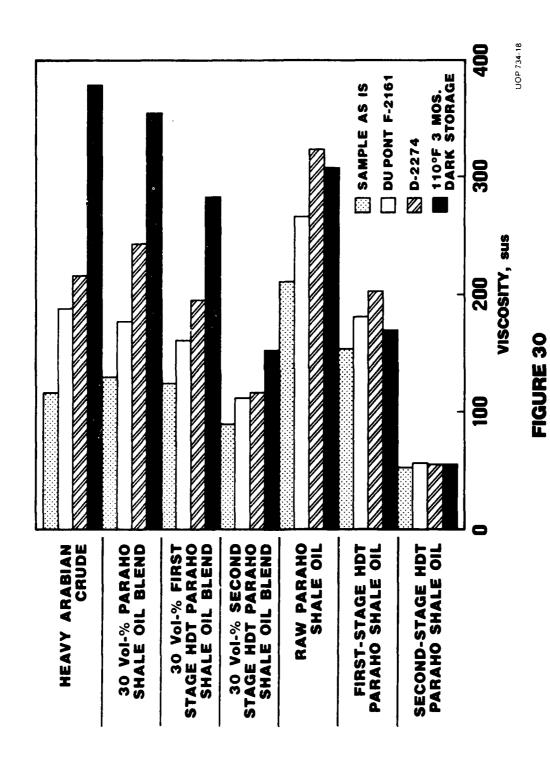
EFFECT OF STORAGE ON HEPTANE INSOLUBLES STUDY SHALE PARAHO

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SECTION VIII

SHALE OIL UPGRADING ECONOMICS

In addition to selecting a novel conversion processing scheme for high jet fuel yield, the proposed flow scheme will provide a realistic and economical solution to the problem of shale oil conversion. The UOP processing scheme, as shown in Figure 31, provides for the removal of arsenic and stabilization of the raw shale oil in a relatively moderate, first-stage pretreatment step. Denitrification is accomplished in a more severe, second-stage hydrotreatment. Finally, the upgraded shale oil is converted to transportation fuels in the hydrocracker. Hydrocracker jet and diesel fuel products do not require further treatment.

Study Basis

The basis for establishing the processing scheme and developing reliable estimates of external requirements and investment costs for the key processing units includes the following:

- A. The refinery design and flow schemes are based on processing 100,000 barrels per stream day (BPSD) of desalted and deashed Occidental shale oil, described in Table 50.
- B. Two primary products -- JP-4 and JP-8 aviation turbine fuels -- are required. Coproducts such as various grades of diesel fuel, fuel oil, motor gasoline, aviation gasoline, or other grades of turbine fuel may be produced.
- C. Only commercially proven processes are to be considered in the refinery processing schemes.
- D. There are no outside feed streams to be processed with the raw shale oil. Fuels for heating are to be internally generated.

E. The refinery is assumed to be a grassroots facility located at a Rocky Mountain site.

F. Capital Investment

- 1. Western U.S. location
- 2. 100,000 BPSD crude shale oil capacity
- 3. First quarter 1981 cost base
- 4. 100% equity financing
- 5. Investment timing over a three year constructs in period
 - a. 25% first year
 - b. 50% second year
 - c. 25% third year
- 6. 10% invesment tax credit
- 7. 45% of plant onsites (not including feed and product storage) will be used to generate plant offsite cost.

G. Working Capital

- 1. 21 days crude storage capacity with 14 days crude inventory
- 2. 14 days product storage capacity with 7 day product inventory
- 3. Crude shale oil valued at \$40.00 per barrel
- 4. Product valued at determined cost
- 5. Debt financed at 15%

H. Capital Return

- 1. 15% DCF rate of return on capital
- 2. 13 years sum of years digits depreciation
- 3. Zero salvage value

I. Operating Basis

- 1. 16 year plant operating life
- 2. 50% operating capacity first year, 100% thereafter
- 90% on-stream factor

- 4. 100,000 BPSD capacity
- 5. All process fuel/heat requirements shall be generated internally from the original shale oil feed.

J. Operating Cost Basis

- 1. Crude Shale 0il -- \$40.000 per barrel
- 2. Cooling Water -- \$0.03 per 1000 gal
- 3. Electricity -- \$0.045 per kWh
- 4. Operators -- \$12.00 per hour
- 5. Helpers -- \$10.50 per hour
- 6. Labor Supervision -- 25% of direct labor
- 7. Overhead -- 100% of direct labor
- 8. Federal and State Taxes -- 50%
- 9. Maintenance, Local Taxes and Insurance -- 4.5% of fixed investment
- 10. Product Values -- all liquid fuel products are of equal value
- 11. By-product Values
 - a. Ammonia at \$155/short ton
 - b. Sulfur at \$105/long ton

K. Miscellaneous

- 1. Use English units
- 2. Mass flow rates
 - a. Barrels per stream day (BPSD)
 - b. Short tons per day (ST/D)
 - c. Standard cubic feet per day (SCFD)

Refinery Processing Scheme

The processing scheme for the production of JP-4 and JP-8 jet fuels, shown in Figures 32 and 33, respectively, are nearly identical except for the size of the units. Naphtha hydrotreating and Platforming units are added to the flow scheme when producing JP-8, and a partial oxidation unit

is added to supplement the hydrogen plant when producing the JP-4 in the diesel fuel case. Various combinations of jet fuel and diesel fuel are produced by utilizing the same flow schemes and by varying the hydrocracking severity to produce the desired product slate. Overall material balances for all cases for JP-4 and JP-8 jet fuels are shown in Table 51.

First-Stage Hydrotreating (UOP RCD Unibon)

The desalted shale oil charge stock is combined with amine scrubbed recycle hydrogen gas plus make-up hydrogen gas (from the hydrogen plant), heated to the desired reaction temperature, by means of exchange and direct fired heaters, and charged to the reactors. The reactor effluent, after heat exchange with the incoming feed, is water washed, condensed, cooled and sent to the product separator. The hydrogen rich off-gas from the separator is amine treated and recycled to the reactor section. Ammonia-rich water is removed from the separator and sent to the sour water stripper. Liquid hydrocarbon product from the separator is not stripped or fractionated but pumped directly into the high-pressure, second-stage hydrotreater.

The first-stage hydrotreater saturates the olefins and diolefins by contacting the feedstock with high purity hydrogen over a catalyst bed. Iron and arsenic are removed along with most of the sulfur compounds. Essentially no arsenic and only small amounts of unsaturates remain; however, a high concentration of nitrogen compounds is still present in the first-stage product and must be removed in the more severe second-stage hydrotreating step.

Second-Stage Hydrotreating

The first-stage separator liquid is combined with the recycle hydrogen-rich gas plus make-up hydrogen (from the hydrogen plant), heated to the desired reactor temperature, by means of exchange and direct fired heaters, and charged to the reactors. The reactor effluent is heat exchanged with the reactor feed, water washed, condensed, cooled and sent to the product separator. The hydrogen-rich gas from the separator is

recycled to the reactor, and the ammonia-rich water is drawn from the separator and treated in the sour water stripper. Liquid hydrocarbons from the separator are pressured directly to the hydrocracking unit (HC Unibon) for further processing.

The second-stage hydrotreating step completes the denitrification of the shale oil to less than 1000 wt-ppm. Hydrocarbon liquid to be used as an environmentally acceptable fuel oil for process heaters is drawn from the high-pressure hydrotreater separator. This material will be stripped, with the light ends going to fuel gas treating and the bottoms liquid used as fuel oil. The balance of the high pressure separator liquid will be used as hydrocracker feedstock. The two-stage hydrotreating yields are given in Table 52.

Hydrocracking (HC Unibon)

Hydrocracking is a highly versatile process for the conversion of a variety of petroleum/hydrocarbon fractions to yield more valuable, lower-boiling products. Along with the molecular weight reduction accomplished in the hydrocracking reactions, there is almost complete conversion of sulfur-, nitrogen- and oxygen-containing compounds, thus yielding products that are mixtures of essentially only paraffins, naphthenes and aromatics. The kerosine products yielded from the HC Unibon unit are of good quality. Both kerosine cuts (JP-4 and JP-8) are nearly sulfur free, low in aromatics, have low freeze points and good smoke points. These hydrocracked kerosines make good jet fuel blending stock.

The HC Unibon unit proposed for this refinery is a parallel-flow hydrocracker designed to process the hydrotreated Occidental shale oil into light ends, light and heavy naphthas and kerosine/diesel products. Several operational modes vary the end points on the kerosine to make the JP-4 or JP-8 jet fuels and the DF-2/DFM diesels. Tables 53 through 56 present the yield estimates and product properties when operating for JP-4 and JP-8 jet fuels including the alternative diesel cases.

The hydrotreated shale oil charge stock is combined with the recycle hydrogen-rich gas plus hydrogen make-up, heated to the desired reactor temperature by means of heat exchange and a direct fired heater, and charged to the reactor section. The reactor section effluent is water washed, cooled and sent to the product separator. The hydrogen rich off-gas from the separator is returned to the reactor section as recycle gas and hydrogen quench.

Ammonia is removed from the product separator with the wash water. The hydrocarbon liquid from the separator is flashed to the low pressure flash drum where much of the H₂S and light gases are removed to the fuel gas treating system. The low-pressure, flash hydrocarbon liquid is charged to a fired heater and sent to the product fractionator where the IBP-300°F material is taken overhead, the 300-520°F kerosine is stripped and blended with the C5+ naphtha product from the overhead material into JP-4 jet fuel. For the JP-8 jet fuel case, a 300-550°F kerosine is stripped for flash point and sent to the product storage. When diesel products are required with the jet fuels, a lower side-cut draw is made on the product fractionator, and a side-cut stripper is provided to make The product fractionator bottoms material is combined with flash point. recycle hydrogen gas and make-up hydrogen and charged to the reactor section. Utilties consumption figures can be found in Tables 62 through 65, and capacities and investments are shown in Table 66.

Light Ends Fractionation

The product fractionator overhead material is cooled, condensed and then collected in the overhead receiver. The overhead receiver gas is compressed and recontacted with the overhead naphtha liquid in the recontact drum. Net gas is removed to fuel gas treating, and the naphtha is sent to the naphtha splitter, where C_6 minus material is taken overhead and the C_7 -300°F heavy naphtha is taken as bottoms product for blending into the JP-4 jet fuel, or used as feed to the naphtha hydrotreater and Platformer to make high octane gasoline when the plant is making JP-8 jet fuel. The naphtha splitter overhead (C_6 -) material is charged to a debutanizer to remove C_4 - material from the light naphtha (C_5/C_6). The

debutanized light naphtha is blended with the heavy naphtha $(C_7-300^{\circ}F)$ and kerosine $(300-520^{\circ}F)$ to make JP-4, the wide-cut jet fuel or the light naphtha is blended into the gasoline pool with high-octane Platformate produced during the JP-8 jet fuel production. If required, part of the light naphtha can be utilized in the hydrogen plant as fuels/feedstock. A small depropanizer is used to separate the C_3 and lighter material from the butanes in the C_4- overhead liquid from the debutanizer. The depropanizer bottoms butanes are used to make vapor pressure in the gasoline during the JP-8 operation. Leftover light ends are sent to fuel gas treating prior to being used as hydrogen plant feedstock. Utilities consumption figures can be found in Tables 62 through 65, and capacities and investments are shown in Table 66.

Naphtha Hydrotreating (For JP-8 Cases Only)

The C7-300°F hydrocracked naphtha from the bottoms of the naphtha splitter is charged to the naphtha hydrotreater, for complete desulfurization and other contaminant removal (oxygen and nitrogen), prior to reforming in a UOP Platforming unit. The naphtha is combined with hydrogen-rich recycle gas plus make-up hydrogen gas, heated to the desired reaction temperature by means of exchange and direct fired heater, and charged to the reactor. The effluent from the reactor, after heat exchange with the incoming feed, is condensed, cooled and sent to the product separator. The hydrogen rich off-gas from the separator is recycled to the reactor section while net separator gas is sent to fuel gas.

The liquid product from the separator is sent to a stripper column for hydrogen sulfide and light ends removal. The C₆ plus stripper bottoms product is charged directly to the Platformer. Table 57 shows the naphtha hydrotreater yield estimate, while Tables 64 and 65 list the utilities. Table 66 presents the plant capacities and investment costs.

UOP Platforming (For JP-8 Cases Only)

In the UOP Platforming unit the hydrotreated naphtha is combined with the recycle hydrogen-rich gas stream, heated by means of exchange and a The hydrotreated shale oil charge stock is combined with the recycle hydrogen-rich gas plus hydrogen make-up, heated to the desired reactor temperature by means of heat exchange and a direct fired heater, and charged to the reactor section. The reactor section effluent is water washed, cooled and sent to the product separator. The hydrogen rich off-gas from the separator is returned to the reactor section as recycle gas and hydrogen quench.

Ammonia is removed from the product separator with the wash water. The hydrocarbon liquid from the separator is flashed to the low pressure flash drum where much of the H2S and light gases are removed to the fuel The low-pressure, flash hydrocarbon liquid is gas treating system. charged to a fired heater and sent to the product fractionator where the IBP-300°F material is taken overhead, the 300-520°F kerosine is stripped and blended with the C5+ naphtha product from the overhead material into JP-4 jet fuel. For the JP-8 jet fuel case, a 300-550°F kerosine is stripped for flash point and sent to the product storage. products are required with the jet fuels, a lower side-cut draw is made on the product fractionator, and a side-cut stripper is provided to make flash point. The product fractionator bottoms material is combined with recycle hydrogen gas and make-up hydrogen and charged to the reactor section. Utilties consumption figures can be found in Tables 62 through 65, and capacities and investments are shown in Table 66.

Light Ends Fractionation

The product fractionator overhead material is cooled, condensed and then collected in the overhead receiver. The overhead receiver gas is compressed and recontacted with the overhead naphtha liquid in the recontact drum. Net gas is removed to fuel gas treating, and the naphtha is sent to the naphtha splitter, where C_6 minus material is taken overhead and the C_7 -300°F heavy naphtha is taken as bottoms product for blending into the JP-4 jet fuel, or used as feed to the naphtha hydrotreater and Platformer to make high octane gasoline when the plant is making JP-8 jet fuel. The naphtha splitter overhead $(C_6$ -) material is charged to a debutanizer to remove C_4 - material from the light naphtha (C_5/C_6) . The

debutanized light naphtha is blended with the heavy naphtha (C7-300°F) and kerosine (300-520°F) to make JP-4, the wide-cut jet fuel or the light naphtha is blended into the gasoline pool with high-octane Platformate produced during the JP-8 jet fuel production. If required, part of the light naphtha can be utilized in the hydrogen plant as fuels/feedstock. A small depropanizer is used to separate the C3 and lighter material from the butanes in the C4- overhead liquid from the debutanizer. The depropanizer bottoms butanes are used to make vapor pressure in the gasoline during the JP-8 operation. Leftover light ends are sent to fuel gas treating prior to being used as hydrogen plant feedstock. Utilities consumption figures can be found in Tables 62 through 65, and capacities and investments are shown in Table 66.

Naphtha Hydrotreating (For JP-8 Cases Only)

The C7-300°F hydrocracked naphtha from the bottoms of the naphtha splitter is charged to the naphtha hydrotreater, for complete desulfurization and other contaminant removal (oxygen and nitrogen), prior to reforming in a UOP Platforming unit. The naphtha is combined with hydrogen-rich recycle gas plus make-up hydrogen gas, heated to the desired reaction temperature by means of exchange and direct fired heater, and charged to the reactor. The effluent from the reactor, after heat exchange with the incoming feed, is condensed, cooled and sent to the product separator. The hydrogen rich off-gas from the separator is recycled to the reactor section while net separator gas is sent to fuel gas.

The liquid product from the separator is sent to a stripper column for hydrogen sulfide and light ends removal. The C_6 plus stripper bottoms product is charged directly to the Platformer. Table 57 shows the naphtha hydrotreater yield estimate, while Tables 64 and 65 list the utilities. Table 66 presents the plant capacities and investment costs.

UOP Platforming (For JP-8 Cases Only)

In the UOP Platforming unit the hydrotreated naphtha is combined with the recycle hydrogen-rich gas stream, heated by means of exchange and a fired heater, then charged successively through the reforming reactors. Fired interheaters are provided between the reactors in order to achieve the desired reactor inlet temperatures.

The effluent from the last reactor, after heat exchange with the incoming feed, is condensed, cooled and sent to the product separator. The hydrogen-rich gas stream from the separator is recycled to the reactor section while the off-gas produced is used as make-up hydrogen for the naphtha hydrotreater and as feedstock to the hydrogen plant. The separator hydrocarbon liquid after recontacting with the compressed separator gas, is separated and charged to the stabilizer (debutanizer for motor fuel operation) where the light hydrocarbons are taken overhead to hydrocracker fractionation and the C5+ Platformate is sent to gasoline blending.

The UOP Platformer is designed for processing C7-300°F naphtha in a conventional fixed bed, semi-regenerative type reformer, featuring low pressure operation with gas recontacting giving maximum liquid recovery and high hydrogen purity. The yields and some of the major properties are noted in Table 58. The units are designed for 100 RON clear Platformate for the maximum JP-8 case, and 98 RON clear Platformate for the JP-8 plus diesel case. The Platformer is designed for conventional catalyst regeneration with an estimated first cycle length of twelve months.

The liquid C₄- material from the Platformer debutanizer is sent to the depropanizer in the fractionation section to recover butanes for gasoline blending. There is no attempt to recover propane, and all the light end hydrocarbons are taken to the fuel gas system for feed to the hydrogen plant. Part of the hydrogen produced from the Platformer net separator gas is used in the naphtha hydrotreater and the balance of the gas, the larger portion, feeds the hydrogen plant. Tables 64 and 65 present the utilities consumption, and Table 66 shows the capacities and investment requirements for the UOP Platforming unit.

Hydrogen Plant

A hydrogen plant is required to supply 97% hydrogen to operate the hydrogen consuming process units. Estimated hydrogen requirements and hydrogen plant sizes and yields are given in Table 59 for JP-4 cases and Table 60 for JP-8 cases.

The preferential feedstock to the hydrogen plant is treated fuel gas but where fuel gas is in short supply, light naphtha (C_5/C_6) is then used as a supplement. In the JP-4 plus diesel case, there is not enough fuel gas or light naphtha available, so a partial oxidation plant is used to obtain the balance of the hydrogen.

The basic steps for the steam-hydrocarbon reforming process in the production of hydrogen are desulfurization, two-stage CO conversion, CO_2 removal, methanation and compression to the required pressure levels.

A variety of feedstocks can be used, such as natural gas, methane, ethane, propane and butanes. Liquid feeds such as light naphthas are sometimes used. The feed should contain less than 0.5% olefins to prevent carbon formation. The hydrocarbons are charged to the charcoal absorber for sulfur removal then sent to the reformer furnace with steam. The gas is converted largely to hydrogen, CO and CO2. The gases are cooled by the addition of steam or condensate and all are passed over catalyst in the CO converter. About 90 to 95% of the CO is converted to CO2. The hot gases leaving the converter are cooled and then scrubbed with an amine solution to remove most of the CO2. The gases are heated up after leaving the CO2 absorber by exchange with gas coming from the CO converter and pass over catalyst in the methanator. Here all the carbon oxides are converted to methane by reaction with hydrogen. The product hydrogen gas from the methanator, saturated with water vapor, is cooled to 100°F.

Tables 62, 63, 64, 65 and 66 list the hydrogen plant utilities requirement, capacities and investment cost for each case.

Partial Oxidation (For JP-4 plus Diesel Case Only)

Even with lower hydrogen consumption in the JP-4 plus diesel case, the reduced supply of light feedstock (fuel gas and light naphtha) indicates that a partial oxidation unit is needed to supplement the hydrogen production from the steam reformer. The partial oxidation process is very flexible in regard to feedstocks. Residual oils from 1 to 10 °API gravity have routinely been processed. Raw shale oil could conceivably be used as feed but, because of the high arsenic content and other contaminants, the resulting hydrogen gas may be of poor quality. To avoid this problem, hydrotreated shale oil is used as the feedstock.

The treated shale oil is vaporized and mixed with oxygen and steam in the reactor. The first reaction phase is that of heating and cracking the hydrocarbon and dispersing it into oxygen (supplied by an air separation plant included in the scheme) and steam which acts as a moderator. In the second phase of reaction, the hydrocarbon burns when it reaches ignition temperature. The residence time is insufficient for complete combustion of the hydrocarbon, consequently one of the by-products is soot. A direct quench cools the gas and scrubs out the soot in a scrubber which is followed by the CO conversion step. The final composition of the product gas containing carbon dioxide and hydrogen is determined by the equilibrium condition of the shift reaction. The shift reaction is followed by carbon dioxide removal and a final methanation step to obtain the high purity hydrogen as a final product.

Estimated yields for the partial oxidation unit are shown in Table 61. Tables 63 and 66 show the partial oxidation unit utilities consumption, capacity and investment cost.

Amine Treating and Sulfur Recovery

Amine treating of refinery gases removes hydrogen sulfide and other acid gas constituents thus avoiding atmospheric pollution by SO_2 from combustion of these gases. The amine treating process is based on an absorption-regeneration cycle using aqueous solutions of an alkanolamine

which reacts with acid gases. Hydrogen sulfide containing feed is contacted counter-currently with amine solution in an absorption or extraction column. Regenerated amine solution is fed into the top of the column and the rich solution taken from the bottoms through exchange to regeneration, where the acid gases are stripped with steam and recovered as hydrogen sulfide with the regenerated solution recycled back to the absorber. The hydrogen sulfide from the amine unit and sour water from the sour water stripper are charged to a Claus sulfur recovery unit for conversion to elemental sulfur.

The low-pressure hydrotreater (RCD Unibon) recycle gas and the net off-gases from the HC Unibon and naphtha hydrotreater are amine treated for hydrogen sulfide removal. The off-gases are treated prior to being used as feedstock in the hydrogen plant or fuel gas to the heaters. In these processing schemes most of the treated light gases are used as hydrogen plant feedstock with little left over for heating.

Sour Water Treating

In the hydrocracking and hydrotreating processing units, water is injected to prevent build-up of salts on the surface of heat exchange tubes. Wash water used to dissolve ammonia and some hydrogen sulfide cannot be disposed of in the conventional refinery sewer system because of its chemical content. This sour water is charged to a stripper where hydrogen sulfide and ammonia are steam stripped from the water. The stripped water is returned to the refinery and the released gases, hydrogen sulfide and ammonia, are separated with the hydrogen sulfide processed in the sulfur plant for elemental sulfur and the ammonia recovered in a liquefaction plant.

Recovered Products

Aviation Turbine Fuels

The refinery is designed to produce two types of aviation turbine fuels as the major products. These are:

A. <u>JP-4 Jet Fuel</u> -- This military jet fuel is wide-cut with a distillation range from C₅-518°F and with gravity limitations between 45-57 °API. Table 67 compares the product qualities of the various cases with the Military Specifications for JP-4. The maximum JP-4 case product meets these specifications, but the JP-4 in the JP-4 plus diesel case has a slightly higher distillation than allowed in the specification. The compositions of the JP-4 jet fuels are:

Component	Мах	Max. JP-4 Case			JP-4 + Diesel Case			
	BPSD	ST/D_	°API	BPSD	ST/D	°API		
Lt. Naph. (C ₅ /C ₆)	11,819	1,375.1	81.4	5,601	649.7	82.1		
C7-300°F Naph.	7,100	947.3	54.2	3,047	406.6	54.2		
300-520°F Kero.	72,841	10,230.7	44.9	45,307	6,363.5	44.9		
TOTAL	91,760	12,553.1	49.6	53,955	7,419.8	48.7		

B. <u>JP-8 Jet Fuel</u> -- This military jet fuel is of the kerosine type, similar to ASTM Jet A-1 fuel, with a maximum distillation end point of 572°F, and with gravity limitations between 37-51 °API. Table 67 shows a comparison between the estimated product for each case with the JP-8 Military Specification. The jet fuel product does meet, or exceed, these specifications. The quantity and quality of the JP-8 jet fuels are:

Component	Max	k. JP-8 Case	е	JP-8	+ Diesel	Case
	BPSD	ST/D	°API	BPSD	ST/D	°API
300-550°F Kero.	80,742	11,380.1	44.3	55,219	7,782.8	44.3

Diesel Fuels

In the alternative cases for JP-4 and JP-8, diesel fuels are produced along with the respective jet fuels. The diesel fuels recovered from these cases satisfy both the DF-2 Diesel and Marine Diesel Specifications. As noted in Table 68, the JP-4 + DF-2/DFM and JP-8 + DF-2/DFM diesels are very similar except for slight differences in properties. The quantity and composition of the diesel fuels are:

Component	JP-4 +	DF-2/DFM	Case	JP-8 +	- DF-2/DFM	Case
	BPSD	ST/D	°API	BPSD	ST/D	°API
520-700°F Diesel	36,871	5,376.9	38.4	-	-	-
550-700°F Diesel		_	_	29,883	4,373.0	37.8

Motor Gasoline Fuels

Motor gasolines are produced only when JP-8 jet fuels are made. The motor gasoline blended in this study is special grade unleaded, which meets Military Specification and Federal Specification VV-G-1690B dated July 1, 1978. Table 69 compares the study blends with the required specification and the blends equal, or exceed, these specifications. The composition of the gasoline blends is as follows:

Max. JP-8 Case Gasoline Blend

Component	BPSD	ST/D	Sp. Gr.	RVP	RON Clear	MON Clear	R+M/2 Clear
Platformate	5,139	740.8	0.8233	4.0	100.0	88.6	
Lt. Naphtha	2,776	322.8	0.6643	11.0	75.0	73.0	
Mixed Butanes	934	93.9	0.5743	_	98.5	94.3	
TOTAL	8,849	1,157.5	0.7471	13.5	92.0	84.3	88.1

JP-8 + DF-2/DFM Case Gasoline Blend

Component	BPSD	ST/D	Sp. Gr.	RVP	RON Clear	MON Clear	R+M/2 Clear
Platformate	2,337	333.8	0.8159	3.9	98.0	87.0	
Hvy. Naphtha	280	37.3	0.7620	0.3	59.0	57.0	
Lt. Naphtha	454	52.7	0.6625	11.0	75.0	73.0	
Mixed Butanes	450	45.3	0.5743		98.5	94.3	
TOTAL	3,521	469.1	0.7609	13.5	92.0	83.7	87.9

It should be noted that both grades of gasoline meet or exceed the octane requirments and the other properties required by the gasoline specs. The composition of these blends is toward a light gasoline with a large percentage of gasoline composed of light naphtha and butanes, but the blends are acceptable by the specifications.

Refinery Fuel Gas

As the material balances shown in Table 51 indicate, no fuel gas is used for heating in any of the processing schemes. All the fuel gas is treated and sent to the hydrogen plant as feedstock.

Refinery Fuel 0il

All heating will be done with fuel oil or naphtha that is not processed into gasoline. Fuel oil is drawn from the refinery processing scheme after the second-stage hydrotreater. The high-pressure separator liquid, drawn for fuel oil, is stabilized in the fuel oil stabilizer with the light hydrocarbon material going to fuel gas treating and the stabilized liquid used as fuel oil. The light naphtha used as fuel in the Max. JP-8 case can be burned in the hydrogen plant. Table 51, Overall Material Balance, indicates the quantities of fuel oil used in each case.

Recovered By-Products

Sulfur -- Approximately 97.5 short tons per stream day of sulfur are recovered from the sulfur plant in each of the processing cases.

Ammonia -- Between 306-307 short tons per stream day of liquid ammonia are recovered from the ammonia liquefaction plant in each of the processing cases.

Operating Cost

The overall refinery operating costs for the JP-4 and JP-8 cases are shown in Tables 70 through 7. Operating costs are divided into direct and indirect operating costs which are as follows:

Direct Operating Cost

Refinery Labor -- Refinery labor includes all the personnel for the process units hired at \$12.00 per hour for operators and \$10.50 per hour

for helpers. To arrive at a total labor cost, a 100% overhead allowance is added to these base rates. A 25% allowance for supervision is also made.

Maintenance Allowance — The maintenance allowance covers normal operating maintenance and turnaround contract maintenance for all refinery equipment including process units, offsites and depreciable assets. An amount equal to 3% of erected plant investment is allocated for maintenance.

<u>Utilities</u> — Refinery fuel is generated internally for both JP-4 and JP-8 cases. Power is purchased at \$0.045 per kWh, and cooling water is priced at \$0.03 per 1000 gallons. Boiler feed water is estimated at \$0.50 per 1000 pounds. Steam (600 psig) is generated at \$0.68 per 1000 pounds (fuel provided from internal sources). Cold treated water is purchased at \$0.07 per 1000 pounds. All utilities are consumed at normal average operating rates.

<u>Catalyst</u>, <u>Solvents and Chemicals</u> — Catalyst consumption is based on the expected catalyst life for normal operating conditions. Similarly, solvents and chemicals are based on normal average operating usage.

Indirect Operating Cost

Local Taxes -- An allowance of 1% of erected plant investment, catalyst and working capital is allocated toward local taxes.

<u>Insurance</u> -- An allowance of 1/2% of erected plant investment, catalyst and working capital is allocated toward insurance.

Process Design, Capital Cost and Utilities

Process designs were prepared for the first- and second-stage hydrotreaters and for the primary conversion section, the hydrocracker. These designs were based on the yields and operating conditions developed from the specific laboratory and pilot plant data generated during the

earlier phases of this project. Each process design includes a heat and material balance, characteristics of principal equipment, and a detailed process flow diagram. The heat and material balance includes stream quantities (mass and volume) and compositions (H_2 , H_2S , NH_3 , H_2O , light hydrocarbons), molecular weights, flowing densities and enthalpies entering and leaving each major equipment item, in addition to pressures and temperatures.

In order to provide more definitive cost data for these primary units, the process design was used as a basis for generating EFCEST (Engineering for Cost Estimating) data. EFCEST data provide a sufficient level of detailed project engineering work to appropriately describe the equipment required. The EFCEST data were then used to prepare a detailed cost estimate with an estimated precision of ±25% for first quarter, 1981 U.S. Gulf Coast construction. To obtain a Western location construction cost, the Gulf Coast cost for plant and offsites were increased by 5%.

Material balances prepared for the naphtha hydrotreating and Platforming units, amine treating and sulfur recovery, sour water stripping, raw feed desalting, ammonia plant and hydrogen production units include stream quantities and compositions. Considering the extent of UOP experience with these types of units, reliable cost and utility estimates were provided without preparing EFCEST data and detailed cost estimates. The material and labor estimates provided for the above units were derived by scaling detailed estimates prepared for similar units.

In Table 66, the capacity and the estimated erected cost (EEC) for individual process units are shown. The costs for the JP-4 and JP-8 cases are similar with the exception that a naphtha hydrotreater and Platformer unit combination is included in the JP-8 case, and a partial oxidation unit is added to the JP-4 plus diesel case. The common facilities cost includes such items as buildings (process control house and substations), electrical power distribution, area lighting and site development in the process area.

Offsite costs are influenced more by the characteristics of the site, local regulations and policies of the refinery than by minor differences in process scheme. Allowances for offsite costs are included, with recognition of special requirements such as for waste treatment, where applicable. The offsite facility cost estimates were based on 45% of the plant onsite investment plus the cost of specified tankage capacity for storing crude and products. This basis was used to estimate offsite facility costs as no other design information was available. As such, the offsite cost is an order-of-magnitude cost estimate and has no meaningful accuracy range.

The capital investment summary for the JP-4 and JP-8 cases is presented in Table 74. The total capital investment is estimated to be \$902 x 10^6 for the maximum JP-4 case and \$939 x 10^6 for the maximum JP-8 case. For the alternative diesel cases, the JP-4 + DF-2/DFM case has a investment of \$989 x 10^6 and the JP-8 + DF-2/DFM case a \$905 x 10^6 investment.

Utility requirements are estimated from the process design data and the information provided for the auxiliary units. Utilities estimated include: electric power, fuel, steam at recommended pressure levels, boiler feedwater, condensate, cold process water and cooling water.

Linear Programming

The use of UOP's linear programming (LP) capabilities permitted the evaluation of numerous processing and product blending alternatives required in this shale oil upgrading study. The linear programming technique, using a multitude of mathematical calculations, allowed various alternatives to be quickly examined to determine the best or optimal processing scheme. Once developed, LP models can be used to perform sensitivity analysis, wherein the effect of varying feedstock value, product prices, quantities and specification can be considered.

The UOP LP system was developed over the years to apply this powerful mathematical method to the analysis of a grassroots refinery and expansion

projects. LP techniques are effectively used in: feedstock evaluation and selections, operations planning, financial planning for new facilities or expansion of existing ones, and development of planned turnaround maintenance. Combining the market data of product prices and demands with data describing technology and economic factors, the LP model can be used to evaluate many alternatives quickly and efficiently. Although the results are a linear representation of reality, they can give the planner a logical basis for a systematic approach to solving these problems.

To illustrate the mathematical relationships and the extent to which the refinery interactions are considered, a matrix representation of the linear equations is used to describe the refinery. In this arrangement, data relationships cascade from one activity to the next, much the same way material flows through the refinery. The model can consider the following general components of refinery profitability:

- Availability of raw materials
- Component blending relationships and recycle streams
- Process unit material balances
- Addition requirements for product quality improvements
- Utility consumptions and conversions (purchase vs. internal generation)
- Market demands for finished products.

The constraints or limitations on the model are contained in the rows of the matrix. The constraints considered in the refinery optimization are:

- Raw material purchase limits
- Unpooled utility limits to sale or purchase
- Product quality specifications
- Material balances on all streams in the refinery
- Capacity limits in process units
- Additive blending
- Utility balances (choice between purchase and internal generation)
- Product demands at market centers
- Physical and social requirements (e.g., emission limits)

Investment in onsite process equipment and related offsite facilities is handled in a manner that considers the non-linear relationship between capital cost and plant capacity. UOP's linear programming system has the ability to recurse on the capital cost to ensure that the capital charges used by the model are consistent with the calculated process unit capacity. The effect of physical and social policy requirements can be evaluated by placing additional constraints on the model and evaluating the results.

The output of the LP program is in the form of reports that allow management to evaluate the results in familiar terms including such terms as:

- Material balances including plant yields and product blends
- Processing schemes
- Utility balances
- Investment requirements
- Operating requirements
- Production costs

In addition, the LP allows the investigation of the sensitivity of the solution through a technique called post optimal analysis. This feature permits the quantified measurement of varying key parameters such as:

- Sales of selected products on various markets
- Capacity of particular process units
- Finished product qualities
- Prices of products and feedstocks

Conclusions

The total production costs for the JP-4 and JP-8 cases are tabulated in Table 75. Incorporating the feedstock cost, operating costs, capital charges, taxes and a 15% DCF rate, and assuming that all liquid products have equal value, the cost of total liquid fuel production in terms of \$/Bbl of shale oil feed are as follows:

Case	\$/Bbl of Feed
Max. JP-4	52.14
JP-4 + DF-2/DFM	53.42
Max. JP-8	52.68
JP-8 + DF-2/DFM	52.15

The difference in cost for the various cases is almost entirely due to the variations in capital cost, especially in the JP-4 + DF-2/DFM case which has the partial oxidation unit.

The product costs can be stated on a "per barrel of total liquid feed" basis and this is calculated by dividing production costs by the volume fraction yield of liquid fuel. Tabulated below are the results of the calculation for each case:

Case	Total Liquid Product, BPSD	\$/Bbl of Liquid Froducts
Max. JP-4	91,760	56.82
JP-4 + DF-2/DFM	90,826	58.82
Max. JP-8	89,591	58.8 0
JP-8 + DF-2/DFM	88,623	58.85

A more detailed analysis of these four cases is provided in the proforma financial statements attached as Appendices B.1 to B.4 and in the Standard Optimization Reports attached as Appendices B.5 to B.8.

TABLE 50. SHALE OIL INSPECTION

Occidental Shale Oil

Feed Source	Desalted Shale Oil
Gravity, °API	22.9
Distillation (D-1160), °F	
IBP	376
50%	712
EP	953
% Over	87
Sulfur, wt-%	0.64
Nitrogen, wt-%	1.51
Pour Point, °F	+65
BS and W, vol-%	0.2
Conradson Carbon, wt-%	1.36
Carbon, wt-%	79.3
Hydrogen, wt-%	12.2
Oxygen, wt-%	0.65
Metals, wt-ppm	
Arsenic	27.5
Iron	23.0
Nickel	10.0
Vanadium	5.0
Ash, wt-%	0.014
Bromine Number	23.5

TABLE 51. OVERALL MATERIAL BALANCES

1

JP-4 and JP-8 Jet Fuel Cases

		Max. JP-4		-d	JP-4 + DF-2/DFM	DFM	,	Max. JP-8		-9L	JP-8 + DF-2/DFM	DFM
Feed	W+-W	No!-8	BPSD	W+-8	Vo!-%	BPSD	W+-8	Vol-8	BPSD	×+-8	%-10A	BPSD
Shale Oil	100,00	100,00	100,000	100,00	100,00	100,000	100.00	100,00	100,000	100.00	100,00	100,000
Liquid Products												
Gasoline Jet Fuel JP-4	78.23	91.76	91,760	46.24	53,96	53,955	7.21	8.85	8,849	2,92	3,52	3,521
Jef Fuel JP-8 Diesel (DF-2/DFM)				33,51	36.87	36,871	70.92	80.74	80,742	48.50	55.22 29.88	55,219 29,883
Total Liquid Products	78.23	91.76	91,760	79.75	90.83	90,826	78.13	89.59	165,68	78.67	88.62	88,623
By-Products												
Sulfur	0.61	ı	ı	0.61	ı	1	0.61	1	1	0.61	ı	1
Ammonia	1.91	•	•	1,91	ı	•	1,92	•	1	1.91	•	1
Water (Net Make)	0.73	0.66	664	0.73	0.66	664	0.73	0.66	664	0.73	0.66	664
Total By-Products	3,25	ı	•	3,25	1	ı	3.26	ı	1	3,25	ı	ı
Streams Utilized as Fuel												
Fuel Oil Naphtha	13.78	14.62	14,622	9.88	10.48	10,481	8,43	8.95	8,947	13,50	14.32	14,319
Totał Fuels	13,78	14.62	14,622	9.88	10.48	10,481	14.00	16.64	16,634	13,50	14.32	14,319
Material Lost in Production	5											
Total Losses	4.74	ı	1	7.12	1	1	4.61	ı	ŧ	4.58	ı	ı

TABLE 52. OVERALL REACTOR YIELDS

Low- and High-Pressure Hydrotreating

	JP-4 a	nd JP-8 Cases
		(SCFB)
Feed	Wt -%_	
Shale 011	100.00	100.00
Hydrogen (Chemical)	2.36	(1425)
Total	102.36	
Products		
Water	0.73	
Ammonia	1.85	
Hydrogen Sulfide	0.68	
Propane and Lighter	0.76	
Butanes	0.32	0.51
Pentanes	0.53	0.78
Hexane +	97.49	103.13
Total	102.36	104.42
Estimated Properties of C ₆	<u>+</u>	
API Gravity	31.8	
Sulfur, wt-%	0.003	
Nitrogen, wt-%	0.09	
C ₇ Insolubles, wt-%	<0.05	
Bromine Number	1.1	
Distillation (D-1160),	°F	
IBP	245	
50%	646	
EP	1034	

TABLE 53. REACTOR YIELDS

HC Unibon

		Max	lmum JP-4	
Feed		Wt -%_		SCFB) LV-%
HT Shale Oil* Hydrogen (Chemical)		100.00		00.00 (885)
Total		101.57		
Products				
Ammonia Propane and Lighter Butanes Light Naphtha (C ₅ /C ₆) Heavy Naphtha (C ₇ -300°F) Kerosine (300-520°F)		0.11 3.53 4.85 11.25 6.98 74.85		14.49 7.83 79.84
Total		101.57	1	02.16
Estimated Properties	Charge	<u> c₅/c</u> 6	Naphtha	Kerosine
Dist. Range, °F API Gravity Sulfur, wt-ppm Nitrogen, wt-ppm	31.8 30 900	81.6 3 < 1	C ₇ -300 54.3 3 < 1	300-520 45.0 3
Distillation (D-1160), 'IBP 50% EP RVP, psi	245 646 1034	90 130 180 12.9	200 240 285	300 404 520
RVP, psi P/N/A, vol-% RON Clear RON + 3 cc TEL/gal Flash Point, °F	- - -	83/15/2 75 93	- 44/49/7 59 78 -	- - - 100
Aromatics, vol-% Freeze Point, °F Naphthalenes, wt-%	- - -	- - -	- - -	9.0 -58 0.4
Constant man		_		2.7

Smoke Point, mm

27

^{*} Total hydrotreater, high-pressure separator liquid.

TABLE 54. REACTOR YIELDS

HC Unibon

	JP-4 + DF-2/DFM		
<u>Feed</u>		(SCFB) LV-%	
HT Shale Oil* Hydrogen (Chemical)	100.00	100.00 (803)	
Total	101.43		
Products			
Ammonia Propane and Lighter Butanes Light Naphtha (C ₅ /C ₆) Heavy Naphtha (C ₇ -300°F) Kerosine (300-520°F) Diesel (520-700°F) Total	0.11 2.02 2.38 5.21 3.09 48.03 40.59	6.76 3.47 51.24 41.67 103.14	
Estimated Properties			

Document of the second	Charge	<u>C5/C</u> 6	Naphtha	Kerosine	<u>Diesel</u>
Dist. Range, °F			C ₇ -300	300-520	520-700
API Gravity	31.8	82.3	54.3	45.0	38.4
Sulfur, wt-ppm	30	3	3	3	3
Nitrogen, wt-ppm	9 00	< 1	< 1	3	3
Distillation (D-1160), °	F				
IBP	245	9 0	200	300	520
50%	646	130	240	404	605
EP	1034	180	285	520	700
RVP, psi	_	12.9	_	-	-
P/N/A, vol-%	_	83/15/2	44/49/7	_	-
RON Clear	_	75	59	-	-
RON + 3 cc TEL/gal	_	93	78	-	_
Flash Point, °F	-	_	_	100	268
Aromatics, vol-%	_	-	-	9.0	-
Freeze Point, °F	_	-	-	-50	_
Naphthalenes, wt-%	_	-	_	0.4	_
Smoke Point, mm	_	-	-	27	-
Pour Point, °F	-	-	-	-	0
Cetane Number	_	-	_	-	56
Cloud Point, °F	_	-	_	-	5
Aniline Point, °F	-	_	-	-	198
Viscosity, cSt at 100°F	-	_	-	-	4.4

^{*} Total hydrotreater, high-pressure separator liquid.

TABLE 55. REACTOR YIELDS

HC Unibon

		Maxi	mum JP-8	
Feed	W	<u>t-%</u>		SCFB) LV-%
HT Shale Oil* Hydrogen (Chemical)		0.00 1.55		00.00 (874)
Total	10	1.55		
Products				
Ammonia Propane and Lighter Butanes Light Naphtha (C5/C6) Heavy Naphtha (C7-300°F)	<u>:</u> •	0.11 3.13 4.25 9.78 5.99	;	12.60 6.73
Kerosine (300-550°F) Total		8.29 1.55	_	33.18 02.51
Estimated Properties	Charge	<u>C₅/C</u> ₆	Naphtha	Kerosin

	Charge	<u>C5/C</u> 6	Naphtha	Kerosine
Dist. Range, °F			C7-300	300-550
API Gravity	31.8	81.7	54.3	44.3
Sulfur, wt-ppm	30	3	3	3
Nitrogen, wt-ppm	900	< 1	< 1	3
Distillation (D-1160),	°F			_
IBP	245	90	200	300
50%	646	130	240	419
EP	1034	180	285	550
RVP, psi	-	12.9	_	_
P/N/A, vol-%		83/15/2	44/49/7	_
RON Clear	-	75	59	_
RON + 3 cc TEL/gal	~	93	78	_
Flash Point, °F	-	_	_	100
Aromatics, vol-%	_	_	_	10
Freeze Point, °F	-	-	_	~58
Naphthalenes, wt-%	-	-	-	1.5
Smoke Point, mm		_		26

^{*} Total hydrotreater, high-pressure separator liquid.

TABLE 56. REACTOR YIELDS

HC Unibon

	JP-8 +	DF-2/DFM
<u>Feed</u>		(SCFB) LV-%
HT Shale Oil* Hydrogen (Chemical)	100.00 1.43	100.00 (803)
Total	101.43	
Products		
Ammonia Propane and Lighter Butanes Light Naphtha (C ₅ /C ₆) Heavy Naphtha (C ₇ -300°F) Kerosine (300-550°F) Diesel (550-700°F) Total	0.11 2.02 2.38 5.21 3.09 56.74 31.88 101.43	6.76 3.47 60.29 32.62 103.14

Esti	mated	Pro	perti	les
------	-------	-----	-------	-----

	Charge	<u>C5/C</u> 6	Naphtha	Kerosine	Diesel
Dist. Range, °F			C ₇ -300	300-550	550-700
API Gravity	31.8	82.3	54.3	44.3	37.8
Sulfur, wt-ppm	30	3	3	3	3
Nitrogen, wt-ppm	900	< 1	< 1	3	3
Distillation (D-1160), °	F				
IBP	245	9 0	200	300	550
50%	646	130	240	419	620
EP	1034	180	285	550	700
RVP, psi	_	12.9	_	_	_
P/N/A, vol-%	_	83/15/2	44/49/7	-	-
RON Clear	-	75	59	_	_
RON + 3 cc TEL/gal	_	93	78	-	-
Flash Point, °F	_	-	-	100	288
Aromatics, vol-%	_	-	-	10	_
Freeze Point, °F	-	-	••	-58	_
Naphthalenes, wt-%	_	_	-	1.5	-
Smoke Point, mm	_	_	-	26	_
Pour Point, °F	_	-	-		0
Cetane Number	-	_	_	_	56
Cloud Point, °F	-	_	_	-	5
Aniline Point, °F	_	-	_	-	200
Viscosity, cSt at 100°F	-	-	-	-	4.5

^{*} Total hydrotreater, high-pressure separator liquid.

TABLE 57. REACTOR YIELDS

Naphtha Hydrotreating Unit

	Max. JP-8 and	JP-8 + DF-2/DFM
Feed	Wt-%_	(SCFB) LV-%
Hydrocracked Naphtha (C7-300°F) Hydrogen (Chemical)	100.00	100.00 (16)
Total	100.03	
Products		
Hydrogen Sulfide Propane and Lighter Butanes Pentanes Hexane + Total	0.02 0.02 0.01 0.02 99.96	100.01 100.01
Properties of Hexane +		
API Gravity Sulfur, wt-% Hydrocarbon Type, vol-%	54.4 0.00005	
P N	44 49	
A	7	

TABLE 58. REACTOR YIELDS

UOP Platforming Unit

	Maximu	ım JP-8	JP-8 + I	DF-2/DFM
Feed	Wt-%	(SCFB) LV-%	Wt-%	(SCFB) LV-%
Hydrotreated Naphtha (C7-300°F)	100.00	100.00	100.00	100.00
Products				
Hydrogen Propane and Lighter	2.65 7.00	(1332)	2.54 6.26	(1276)
Butanes	3.95	5.23	3.53	4.67
C ₅ + Platformate	86.40	79.97	87.67	81.85
Total	100.00		100.00	
Properties of C ₅ + Platformate				
API Gravity	40.4		41.9	
RON Clear	100.0		98.0	
RON + 3 cc TEL/USG	105.2		103.7	
MON Clear	88.6		87.0	
MON + 3 cc TEL/USG	92.7		91.5	
RVP, psi	4.0		3.9	

TABLE 59. ESTIMATED YIELDS Hydrogen Plant -- Steam Reforming

		J	JP-4 Case	Se			JP-4 +	DF-2/	JP-4 + DF-2/DFM Case	
Feed to H ₂ Plant	BPSD	Sp.Gr.	MM	MMSCFD	Wt-%	BPSD	Sp.Gr.	MM	MMSCFD	Wt-%
Treated Fuel Gas Light Naphtha	652 (0.6645	24.3	45.6	95.1	1	r	18.2	35.1	100.0
Total Feed					100.0					100.0
Products										
Hydrogen $\mathtt{CH}_{oldsymbol{\mathcal{L}}}$				247.5	42.7				141.4	44.5
Total H_2 (97% Purity)				255.1	53.2				145.8	55.5
Process Loss					46.8					44.5
Total Products					100.0					100.0
Estimated Process Units H ₂ Re	H ₂ Requirements	nts								
Unit				H2 MMSCFD					H ₂	
Low- and High-Pressure Hydrotreater	reater			154.3					154.3	
Hydrocracker (HC Unibon)				93.2					82.9	
Total Hydrogen Required				247.5					237.2	

0

Hydrogen Required from Partial Oxidation

TABLE 60. ESTIMATED YIELDS

Hydrogen Plant -- Steam Reforming

		5	JP-8 Case	se			JP-8 +	DF-2/	JP-8 + DF-2/DFM Case	
Feed to H ₂ Plant	BPSD	Sp.Gr.	MM	MMSCFD	Wt -%	BPSD	Sp.Gr.	MM.	MMSCFD	Wt-%
Treated Fuel Gas			22.3	43.7	83.9			17.8	35.4	55.8
Light Naphtha	1060	0.6643	· 1	0 1	8.0	5345	0.6625	. 1	Ç ı	41.6
Total Feed					100.0					100.0
Products										
Hydrogen CH_4				252.5	43.7				240.3	42.9
Total H ₂ (97% Purity)				260.3	54.5				247.7	53.4
Process Loss					45.5					46.6
Total Products					100.0					100.0
Estimated Process Units H ₂ Re	H ₂ Requirements	nents								
Unit				H2 MMSCFD					H ₂ MMSCFD	
Low- and High-Pressure Hydrotreater	reate	t u		154.3					154.3	
Hydrocracker (HC Unibon)				98.0					85.8	
Naphtha Hydrotreater				0.2					0.2	
Total Hydrogen Required				252.5					240.3	

TABLE 61. ESTIMATED YIELDS

Hydrogen Plant -- Partial Oxidation

		JP-4 + D	F-2/DFM Case	2
Feed to H2 Plant	BPSD	Sp.Gr.	MMSCFD	Wt -%
HT Shale Oil	6906	0.8640		100.0
Products				
Hydrogen			95.8	24.4
CH ₄			3.0	6.0
Total H ₂ (97% Purity)			98.8	30.4
Process Loss				69.6
Total Products				100.0

TABLE 62. UTILITY CONSUMPTION

Maximum JP-4

	Power, kWh	St	Steam, M 1b/hr	b/hr LP	BFW and Condensate, M 1b/hr	Cooling Water, gpm	Fuel Fired, MM btu/hr	Cold Treated Water, M 1b/hr
Feed Preparation	450	1	1	1	ı	ı	ı	ı
LP Hydrotreating (RCD Unibon)	7,945	-80.2	ı	1	+80.2	112	91	39.4
HP Hydrotreating	5,436	-67.7	1	1	+67.7	1	207	135.2
HC Unibon - Hydrocracking	27,797	-29.5	ı	+12.1	-14.4	2597	677	35.8
Hydrogen Plant (Steam Reforming)	8,045	+177.4	t	l	-576.4	472	1963	ı
Fuel Gas Treating	185	ı	-7.5	-42.4	6*67+	2455	t	ı
Sulfur Plant	100	-1.0	+34.5	6.44+	-83.0	ı	Į	1
Sour Water Treating	351	ı	-213.7	I	+213.7	4141	ı	t
Fractionation	995	ı	t	-38.2	+38.2	733	74	t
Fuel Oil Stabilizer	105	•	1	1	ı	214	31	1
TOTAL	50,980	-1.0	-186.7	-23.6	-224.1	10,724	3043	210.4

NOTES:

Negative sign indicates consumption Positive sign indicates production Maximum air cooling

Fuel fired heating based on net heating value.

Accessed the same of the same

TABLE 63. UTILITY CONSUMPTION

JP-4 + DF-2/DFM

	Power, kWh	St.	Steam, M lb/hr	b/hr LP	BFW and Condensate, M lb/hr	Cooling Water,	Fuel Fired, MM btu/hr	Cold Treated Water, M 1b/hr
Feed Preparation	450	ı	ı	1	ı	1	ı	1
LP Hydrotreating (RCD Unibon)	7,945	-80.2	ı	1	+80.2	112	91	39.4
HP Hydrotreating	5,436	-67.7	ŧ	1	+67.7	ı	207	135.2
HC Unibon - Hydrocracking	23,171	ı	+70.4	+44.3	-148.2	1661	653	34.7
Hydrogen Plant (Steam Reforming)	4,661	+97.5	1	1	-322.9	282	1110	ı
Hydrogen Plant (Partial Oxidation)	27,384	+186,1	1	1	-166.2	21,657	187	ı
Fuel Gas Treating	185	i	-7.5	-42.4	6.67+	2454	I	ı
Sulfur Plant	100	-1.0	+34.5	+44.8	-83.0	1	i	ı
Sour Water Treating	349	ı	-212.4	ı	+212.4	4126	ı	ı
Fractionation	262	1	t	-17.9	+17.9	339	34	1
Fuel Oil Stabilizer	125	•	1	ı	1	254	37	1
TOTAL	70,068	+134.7	-115.0	+28.8	-292.2	30,885	2319	209.3

NOTES:

Negative sign indicates consumption Positive sign indicates production Maximum air cooling Fuel fired heating based on net heating value. 1. 3.

TABLE 64. UTILITY CONSUMPTION

Maximum JP-8

	Power, kWh	Ste	Steam, M 1b/hr	b/hr LP	BFW and Condensate, M lb/hr	Cooling Water, gpm	Fuel Fired, MM btu/hr	Cold Treated Water, M 1b/hr
Feed Preparation	450	1	1	1	ı	ı	ı	ı
LP Hydrotreating (RCD Unibon)	7,945	-80.2	ı	ı	+80.2	112	91	39.4
HP Hydrotreating	5,436	-67.7	ı	1	+67.7	ŧ	207	135.2
HC Unibon - Hydrocracking	29,559	-31.3	ŧ	+12.9	-15.3	2761	720	38.1
Naphtha Hydrotreating	191	ı	ı	1	ı	53	5	ı
Platforming	629	+16.3	-0.2	ı	-17.4	382	87	t
Hydrogen Plant (Steam Reforming)	8,142	+185.1	ī	1	-595.1	468	2016	ı
Fuel Gas Treating	185	1	-7.5	-42.4	6*65+	2454	ı	ı
Sulfur Plant	100	-1.0	+34.5	+44.8	-83.0	ı	ı	ı
Sour Water Treating	358	1	-218.5	ı	+218.5	4188	t	ı
Fractionation	531	i	1	-37.0	+37.0	760	89	ı
Fuel Oil Stabilizer	64	1	1	1	1	131	19	1
TOTAL	53,620	+21.2	-191.7	-21.7	-257.5	11,309	3213	212.7

NOTES:

- 1. 2. 4.
- Negative sign indicates consumption
 Positive sign indicates production
 Maximum air cooling
 Fuel fired heating based on net heating value.

TABLE 65. UTILITY CONSUMPTION

JP-8 + DF-2/DFM

Cold Treated Water, M 1b/hr	l	39.4	135.2	35.9	ı	ı	l	ı	1	ľ	1	ı	210.5
Fuel Fired, MM btu/hr	ı	91	207	9/9	2	37	2057	ı	1	ı	35	31	3136
Cooling Water, gpm	1	112	i	1720	24	160	295	2454	1	4150	393	209	9517
BFW and Condensate, M 1b/hr	1	+80.2	+67.7	-153.4	J	-7.7	-642.6	6.64+	-83.0	+214.8	+19.3	,	-454.8
b/hr LP	ı	1	ı	+45.9	ı	ı	t	-42.4	+44.8	ı	-19.3	1	+29.0
Steam, M 1b/hr	1	ı	1	+72.8	ı	-0-1	ı	-7.5	+34.5	-214.8	ı	1	-115.1
Ste	ı	-80.2	-67.7	ı	i	+7.2	+221.8	I	-1.0	ı	1	•	+80.1
Power, kWh	450	7,945	5,436	23,991	85	279	7,021	185	100	352	274	103	46,221
	Feed Preparation	LP Hydrotreating (RCD Unibon)	HP Hydrotreating	HC Unibon - Hydrocracking	Naphtha Hydrotreating	Platforming	Hydrogen Plant (Steam Reforming)	Fuel Gas Treating	Sulfur Plant	Sour Water Treating	Fractionation	Fuel 011 Stabilizer	TOTAL

NOTES:

Negative sign indicates consumption Positive sign indicates production Maximum air cooling Fuel fired heating based on net heating value. 1. 3.

TABLE 66. PROCESS UNITS CAPACITIES AND CAPITAL INVESTMENTS

	Max. JP-4	• 4	JP-4 + DF-2/DFM	+ Q	Max.	• m	JP-8 + DF-2/DFM	- L
	(S Ton/SD) BPSD	MM Dollars						
Feed Preparation	100,000	6	100,000	σ	100,000	6	100,000	σ
LP Hydrotreating (RCD Unibon)	100,000	59	100,000	59	100,000	59	100,000	59
HP Hydrotreating	102,920	76	102,920	76	102,920	76	102,920	76
Hydrocracking (HC Unibon)	94,000	169	91,095	165	66,965	176	94,320	169
Fractionation	1	9	1	4	ı	۲	1	'n
Fuel Gas Treating	ı	4	ı	4	ı	4	ı	4
Sulfur Plant	(61.5)	v	(97.5)	ø	(61.5)	v	(97.5)	vo
Hydrogen Plant (Steam Reforming)	(819,2)	125	(468.2)	79	(835.9)	127	(795.4)	122
Hydrogen Plant (Partial Oxidation)	ı	•	(317.1)	114	1	1	1	1
Naphtha Hydrotreating	ı	ı	•	1	6,485	4	2,875	2
Platforming	1	•	•	1	6,485	Ξ	2,875	9
Sour Water Treating	19,480	0	19,365	01	19,920	10	19,580	01
Fuel Oil Stabilizer	15,360	٤	18,265	ю	9,400	2	15,040	М
Common Facilities	1	=	t	=	1	=		=
TOTAL PROCESS INVESTMENT		499		561		523		503

Note: Capital Investment as of First Quarter 1981.

TABLE 67. PRODUCT QUALITIES

JP-4 and JP-8 38% Fuels

	JP-4 Military Specs.	Max. JP-4 Case	JP-4 + DF-2/DFM Case	JP-8 Military Specs.	Max. JP-8 Case	JP-8 + DF-2/DFM Case
API Gravity,	45-57	49.6	48.7	37-51	44.3	44.3
Aromatics, vol-\$	25 (max)	7.9	8.2	25 (max)	10.0	10.0
Flash Point, °F	ı	•	•	100 (min)	^ 100	^ 100
freeze Point, °f	-72 (max)	< -72	< - 72	-58 (max)	< -58	< -58
Smoke Point, mm	20 (min)	> 20	> 20	20 (min)	56	26
Sulfur, wt-%	0.40 (max)	0.0003	0.0003	0.40 (max)	0.0003	•0003
Naphthelenes, vol-%	ı	1	•	3.0 (max)	1.5	1.5
Olefins, vol−\$	5.0 (max)	r I	Ę	5.0 (max)	- 1 L	114
Distillation, *F						
18P	Report	108	11	Report	300	300
108	Report	179	200	401 (max)	337	337
20%	293 (max)	250	270	Report	361	361
50\$	374 (max)	374	381	Report	419	419
\$ 06	473 (max)	473	479	Report	512	512
EP	518 (max)	518	520	572 (max)	550	550

TABLE 68. DF-2 and DFM DIESEL FUELS

	DF-2 Diesel Fuel (Oconus) Specs.	Diesel Fuel Marine Specs.	JP-4 + DF-2/DFM Case	JP-8 + DF-2/DFM Case
Gravity, °API	32.9-41.0	Report	38.4	37.8
Flash Point, °F	133 (Min.)	140 (Min.)	268	288
Cloud Point, °F	9 (Max.)	30 (Max.)	5	5
Pour Point, °F	0 (Max.)	20 (Max.)	0	0
Sulfur, wt-%	0.7 (Max.)	1.0 (Max.)	0.0003	0.0003
Cetane Number	45 (Min.)	-	56	56
Aniline Point, °F	-	Report	198	200
Viscosity: cSt at 100°F	1.8-9.5	1.8-4.5	4.4	4.5
Distillation, °F				
50% 9 0%	Report 675 (Max.)	- 675 (Max.)	605 673	620 675
EP	700 (Max.)	725 (Max.)	700	700

TABLE 69. GASOLINE BLENDS AND QUALITIES

Special Grade -- Unleaded

		Max. JP	-8 Case	JP-8 +	Diesel
Gasoline Component		Gaso	line	Gaso	oline
		vo1-%	BPSD	vo1-%	BPSD
					
Platformate		58.1	5,139	66.4	2,337
Heavy Naphtha		_	-	7.9	280
Light Gasoline		31.4	2,776	12.9	454
Mixed Butanes		10.5	934	12.8	450
TOTAL		100.0	8,849	100.0	3,521
Properties	W4 1 4 h a mar	Wa	ID 0		
of Gasoline	Military		JP-8	JP-8 +	
or Gasorine	Specs.	_Gas	soline	Gasc	line
API Gravity	_	9	57.9	54	.5
RON Clear	92.0 (Min.)	g	92.0		2.0
MON Clear	82.0 (Min.)	8	34.3		3.7
R+M/2	87.0 (Min.)		38.1		.9
RVP, psi	13.5 (Max.)		3.5		3.5
Sulfur, wt-%	0.10 (Max.)		0003	0.00	
Aromatics, vol-%	55.0 (Max.)		2.8		.3
Distillation (D-1160)	. °F				
IBP	,		53		84
10%	131 (Max.)		105	1	29
30%	_		145		.71
	n.)-235 (Max.)		183		:71
70%			245		49
90%	365 (Max.)		292	_	
EP	437 (Max.)		342		84
	TOI (FIG.)		342	3	35

TABLE 70. MAX. JP-4 CASE -- ESTIMATED OPERATING COST

Basis: 100,000 BPSD at 90% Operating Efficiency

Direct Operating Cost

Α.	Refinery Labor	MM\$/Yr.	MM\$/Yr.
	Direct Labor		
	76 operators	2.67	
	28 helpers	0.85	
	20 helpels	0.03	
	Total Direct Labor	3.52	
	Supervision		
	25% of Direct Labor	0.88	
	Total Operating Labor	4.40	
	Overhead		
	100% of Operating Labor	4.40	
	Total Labor	8.80	8.80
В.	Maintenance		
	Maintenance Allowance at 3%		22.02
	of Plant Investment		22.92
С.	Utilities		
	D 50 001 117 + 00 0/5/117	10 00	
	Power: 50,981 kWh at \$0.045/kWh	18.09	
	600 1b Steam: 211.3 M 1b/hr at \$0.68/M 1b (cost)	1.13	
	BF Water: 224.1 M 1b/hr at \$0.50/M 1b	0.88	
	Cold Treated Water: 233.8 M 1b/hr at \$0.07/M 1b	0.13	
	Cooling Water: 643.4 M gal/hr at \$0.03/M gal	0.15	
	Total Utilities	20.38	20.38
Ð.	Catalyst Replacement, Solvents and Chemicals		
	First-Stage Mudretreating (DCD Uniber)	3.76	
	First-Stage Hydrotreating (RCD Unibon)		
	Second-Stage Hydrotreating	1.41	
	Hydrocracking (HC Unibon)	2.26	
	Hydrogen Plant	1.35	
			_
	Total Catalysts, Solvents and Chemicals	8.78	8.78
	Total Direct Operating Cost		60.88

TABLE 70. MAX. JP-4 CASE -- ESTIMATED OPERATING COST (Cont'd)

	MM\$/Yr.
Indirect Operating Cost	
Local Taxes	
Local Taxes Allowance at 1% of Plant Investment, Catalyst and Working Capital	8.83
Insurance	
Insurance Allowance at 1/2% of Plant Investment, Catalyst	
and Working Capital	4.41
Total Indirect Operating Cost	13.24
Total Operating Cost	74.12

TABLE 71. JP-4 + DIESEL CASE -- ESTIMATED OPERATING COST

Basis: 100,000 BPSD at 90% Operating Efficiency

Direct Operating Cost

Α.	Refinery Labor	MM\$/Yr.	MM\$/Yr.
	Direct Labor 86 operators 32 helpers	3.00 0.99	
	Total Direct Labor	3.99	
	Supervision		
	25% of Direct Labor	1.00	
	Total Operating Labor	4.99	
	Overhead		
	100% of Operating Labor	4.99	
	Total Labor	9.98	9.98
В.	Maintenance		
	Maintenance Allowance at 3% of Plant Investment		25.62
С.	Utilities		
	Power: 70,069 kWh at \$0.045/kWh 600 lb Steam: None BF Water: 292.1 M lb/hr at \$0.50/M lb Cold Treated Water: 232.5 M lb/hr at \$0.07/M lb Cooling Water: 1853.1 M gal/hr at \$0.03/M gal	24.86 - 1.15 0.13 0.44	
	Total Utilities	26.58	26.58
{) .	Catalyst Replacement, Solvents and Chemicals		
	First-Stage Hydrotreating (RCD Unibon) Second-Stage Hydrotreating Hydrocracking (HC Unibon) Hydrogen Plant	3.76 1.41 1.73 1.21	
	Total Catalysts, Solvents and Chemicals	8.11	8.11
	tel Direct Operating Cost		70.29

TABLE 71. JP-4 + DIESEL CASE -- ESTIMATED OPERATING COST (Cont'd)

	MM\$/Yr.
Indirect Operating Cost	
Local Taxes	
Local Taxes Allowance at 1% of Plant Investment, Catalyst and Working Capital	9.70
Insurance	
Insurance Allowance at 1/2% of Plant Investment, Catalyst	
and Working Capital	4.85
Total Indirect Operating Cost	14.55
Total Operating Cost	84.84

TABLE 72. MAX. JP-8 CASE -- ESTIMATED OPERATING COST

Basis: 100,000 BPSD at 90% Operating Efficiency

Direct Operating Cost

Α.	Refinery Labor	MM\$/Yr.	MM\$/Yr.
	Direct Labor 88 operators 32 helpers	3.07 0.99	
	Total Direct Labor	4.06	
	Supervis on		
	25% of Direct Labor	1.02	
	Total Operating Labor	5.08	
	Overhead		
	100% of Operating Labor	5.08	
	Total Labor	10.16	10.16
В•	Maintenance		
	Maintenance Allowance at 3% of Plant Investment		23.97
С.	Utilities		
	Power: 53,622 kWh at \$0.045/kWh 600 lb Steam: 192.2 M lb/hr at \$0.68/M lb (cost) BF Water: 257.5 M lb/hr at \$0.50/M lb Cold Treated Water: 235.1 M lb/hr at \$0.07/M lb Cooling Water: 678.6 M gal/hr at \$0.03/M gal	19.02 1.03 1.02 0.13 0.16	
	Total Utilities	21.36	21.36
D •	Catalyst Replacement, Solvents and Chemicals		
	First-Stage Hydrotreating (RCD Unibon) Second-Stage Hydrotreating Hydrocracking (HC Unibon) Hydrogen Plant Naphtha Hydrotreating and Platformer	3.76 1.41 2.30 1.38 0.13	
	Total Catalysts, Solvents and Chemicals	8.98	8.98
	Total Direct Operating Cost		64.47

TABLE 72. MAX. JP-8 CASE -- ESTIMATED OPERATING COST (Cont'd)

	MM\$/Yr.
Indirect Operating Cost	
Local Taxes	
Local Taxes Allowance at 1% of Plant Investment, Catalyst and Working Capital	9.20
Insurance	
Insurance Allowance at 1/2% of Plant Investment, Catalyst and Working Capital	4.60
Total Indirect Operating Cost	13.80
Total Operating Cost	78.27

TABLE 73. JP-8 + DIESEL CASE -- ESTIMATED OPERATING COST

Basis: 100,000 BPSD at 90% Operating Efficiency

Direct Operating Cost

Α.	Refinery Labor	MM\$/Yr.	MM\$/Yr.
	Direct Labor		
	88 operators	3.07	
	32 helpers	0.99	
	Total Direct Labor	4.06	
	Supervision		
	25% of Direct Labor	1.02	
	Total Operating Labor	5.08	
	Overhead		
	100% of Operating Labor	5.08	
	Total Labor	10.16	10.16
В.	Maintenance		
	Maintenance Allowance at 3%		
	of Plant Investment		23.10
С.	Utilities		
	Power: 46,222 kWh at \$0.045/kWh	16.40	
	600 lb Steam: 34.9 M lb/hr at \$0.68/M lb (cost)	0.19	
	BF Water: 454.8 M lb/hr at \$0.50/M lb	1.79	
	Cold Treated Water: 235.1 M 1b/hr at \$0.07/M 1b	0.13	
	Cooling Water: 571.0 M gal/hr at \$0.03/M gal	0.13	
	Total Utilities	18.64	18.64
D •	Catalyst Replacement, Solvents and Chemicals		
	First-Stage Hydrotreating (RCD Unibon)	3.76	
	Second-Stage Hydrotreating	1.41	
	Hydrocracking (HC Unibon)	1.79	
	Hydrogen Plant	1.36	
	Naphtha Hydrotreater and Platformer	0.05	
	Total Catalysts, Solvents and Chemicals	8.37	8.37
	Total Direct Operating Cost		60.27

TABLE 73. JP-8 + DIESEL CASE -- ESTIMATED OPERATING COST (Cont'd)

	MM\$/Yr.
Indirect Operating Cost	
Local Taxes	
Local Taxes Allowance at 1% of Plant Investment, Catalyst and Working Capital	8.86
Insurance	
Insurance Allowance at 1/2% of Plant Investment, Catalyst and Working Capital	4.43
Total Indirect Operating Cost	13.29
Total Operating Cost	73.56

TABLE 74. CAPITAL INVESTMENT SUMMARY

(Millions of Dollars)

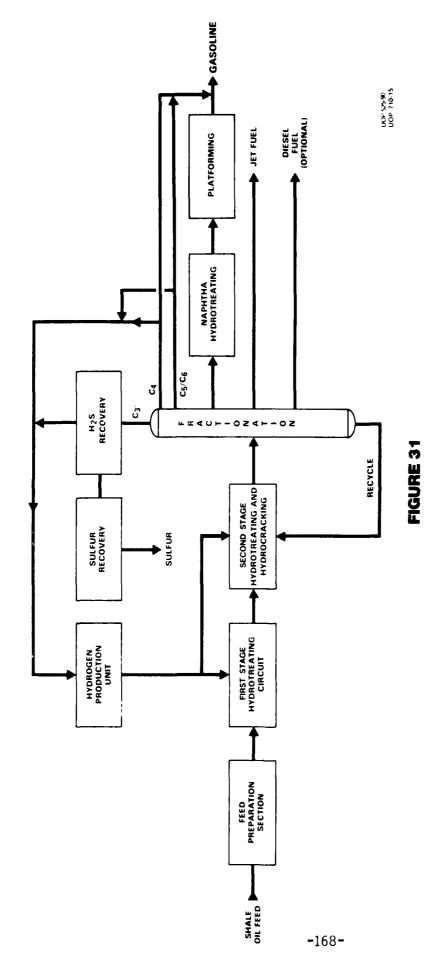
100,000 BPSD Refinery

	Max. JP-4	JP-4 + Diesel	Max. JP-8	JP-8 + Diesel
Process Units Erected Cost	499	561	523	503
Allowance for Offsites	265	293	276	267
Allowance for Fully Paid Royalties and Know-How Fees TOTAL DEPRECIABLE INVESTMENT		<u>20</u> 874	20 819	<u>20</u> 790
Initial Catalyst Inventory	25	22	27	23
Working Capital Allowance	93	_93	93	_92
TOTAL CAPITAL INVESTMENT	902	989	939	905

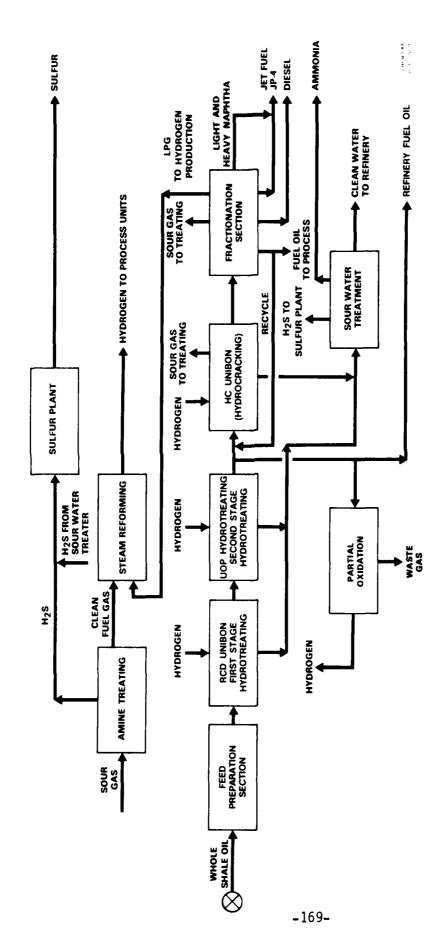
TABLE 75. COST OF PRODUCTION BREAKDOWN

100,000 BPSD Charge Rate

	Max. JP-4	JP-4 + DF-2/DFM	Max. JP-8	JP-8 + DF-2/DFM
Operating Cost, \$/Bb1 of Feed	2.26	2.58	2.38	2.24
Cost of Feed, \$/Bbl	40.00	40.00	40.00	40.00
Capital Charges for 15% DCF Return, \$/Bbl of Feed	9.88	10.84	10.30	9.91
Total Cost of Production, \$/Bbl of Feed	52.14	53.42	52.68	52.15
Total Cost of Liquid Products, \$/Bbl	56.82	58.82	58.80	58.85



UOP SHALE OIL TO FUELS OVERALL BLOCK FLOW DIAGRAM



FRODUCTION OF JP-4 JET FUEL BLOCK FLOW DIAGRAM

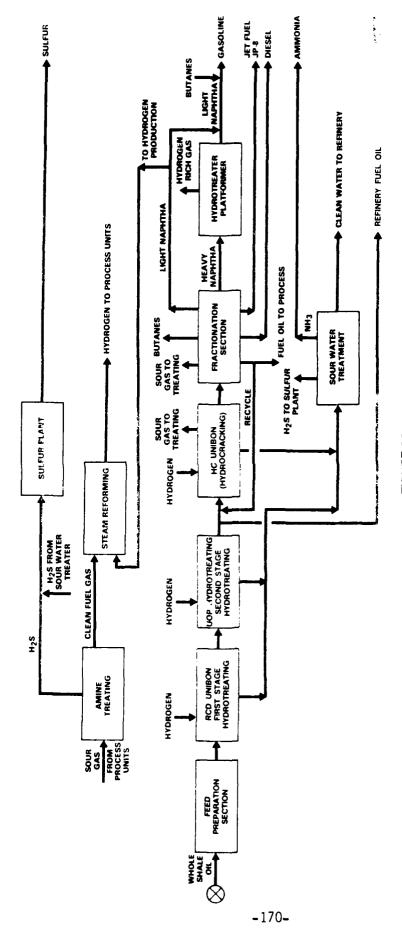


FIGURE 33
PRODUCTION OF JP-8 JET FUEL
BLOCK FLOW DIAGRAM

APPENDIX A

SAMPLE PREPARATION AND ANALYSIS FOR STABILITY/COMPATIBILITY STUDY

A.1 Preparation of Samples

A.1.1 Shale Oil Samples

- 1. Transfer 1850 cc of representative sample to a one-gallon bottle.
- 2. Place one-gallon bottle containing sample in sonic bath. Sonify at $80^{\circ}F$ while bubbling in N_2 at 5 LHSV for 30 minutes to disperse and deoxygenate sample. Seal under N_2 .
- 3. Send samples for the following analysis stored under N_2 .

	Sample Size, g
Distillation (D-1160)	220
Micro C, H, O	1
Sulfur	1
Total N	1
Conradson Carbon	1
C7 Insolubles	1
Toluene Insolubles	1
Viscosity, Kinematic at 100°F	40
Steam Jet Gum	100
Bromine Number	10
Metals by AAS: As, Fe, N_1 , V	25
F 21-51 (300°F Accelerated Fuel Oil	
Stability Test)	60 (mL)
ASTM D 2274-74 (Oxidation Stability Test)	375
Three Months 110°F Dark Storage Scability	375

A.1.2 Petroleum Crude Oil Sample

- 1. To each of 3 one-gallon bottles, transfer 1850 cc of representative sample of petroleum crude oil.
- 2. Sonic disperse and deoxygenate sample as listed in A.1.1.2.
- 3. Send sample for analysis listed in A.1.1.3.

A.2 Preparation of Shale Oil/Petroleum Crude Oil Blends

- 1. Blends are prepared from sonic dispersed and deoxygenated samples. Transfer 300 cc \pm 1 cc shale oil to 1 gallon bottle. Add 700 cc \pm 1 cc petroleum crude oil.
- 2. Place sample blend in sonic bath. Sonify at $80^{\circ}F$ while bubbling in N_2 at 5 LHSV for 5 minutes to disperse and deoxygenate sample. Seal under N_2 .
- 3. Send samples for the following analysis stored under N_2 .

	Sample Si	ze, g
C ₇ Insolubles	1	
Toluene Insolubles	1	
Viscosity, Kinematic at 100°F	40	
Steam Jet Gum	100	
F 21-51 (350°F Accelerated Fuel Oil		
Stability Test)	60	(mL)
ASTM D 2274-74 (Oxidation Stability		
Accelerated Method)	375	
Three Months 110°F Dark Storage Stability	375	

A.3 Three Months 110°F Dark Storage Fuel Oil Stability Test

1. Scope

1.1 This method covers the measurement of the stability of distillate fuel. The fuel is stored in vented Pyrex bottles for three months at 110°F to give an indication of the fuels potential long-term storage stability when stored at lower temperatures.

2. Applicable Documents

2.1 ASTM Standards:

- D 270 Sampling Petroleum and Petroleum Products
- D 2274 Test for Oxidation Stability of Distillate Fuel Oil
 Accelerated Method
- D 1500 ASTM Color of Petroleum Products
- D 381 ASTM Jet Evaporation, Adherent Gums

3. Summary of Method

3.1 A measured volume of filtered fuel is stored in vented Pyrex bottles for three months at 110°F. After three months he samples are examined for color and sediment.

4. Significance

4.1 This test is the most commonly used accelerated test for distillate fuel oils. Many studies have been made and it has been shown that 13 weeks at 110°F dark storage is equivalent, in most cases, to one year of ambient storage (12.1).

5. Apparatus

5.1 Sample containers are Serum Pyrex Bottles (12.2). The total capacity of the container is 500 mL.

F	ÁD-A13	0 774	lica	E CUQI	E OTI	TO FU	FIS DI	IIME 2	PHASE	5 7 AN	D4(II)	IINP	3/4	
ľ			INC JUL	DES F	LAINES WAL-TI	TL P	ELS DL ROCESS 116-VO	DIV L-2 F3	J G S 3615-	IKÓNÏÄ 78-C-2	ĚŤÁL 079	1/4	NL	
	UNCLAS	21+1Ff	<u> </u>								-70 2	1/4		
į														
											_			

- 5.2 A Miracle Top Box (12.3) with cover is used to store samples. Ends are open to allow ventilation.
- 5.3 A hot room is used to store samples. Temperature is thermostatically controlled to maintain a temperature of $110^{\circ} \pm 1^{\circ}$ F.
- 5.4 Drying oven, maintained at $194^{\circ} \pm 2^{\circ}F$.
- 5.5 Filter holder, Millipore, Hydrosol Stainless, 47 mm, XX20 047 20.
- 5.6 Flash 21 filtering with side tube.
- 5.7 Petri dish.
- 6. Reagents and Materials
- 6.1 Filter, 0.8 μ Millipore Type AA.
- 6.2 Hydrocarbon solvent, \underline{n} -heptane, 99% minimum purity, (Phillips Petroleum Company, or equivalent).
- 6.3 All equipment required to determine color by ASTM Method D 1500 (if desired).

7. Sampling Procedure

7.1 A sample for testing will be procured by the method outlined in ASTM Method D 270. Sample containers should preferably be 1-gallon or larger metal can. Sample containers should be rinsed with Adherent Insoluble Solvent (equal parts of reagent grade toluene, acetone, methanol), dried and then nitrogen filled. Samples should be stored at reduced temperature, 33° to 45°F, prior to use.

A.3 Three Months 110°F Dark Storage Fuel Oil Stability Test

1. Scope

1.1 This method covers the measurement of the stability of distillate fuel. The fuel is stored in vented Pyrex bottles for three months at 110°F to give an indication of the fuels potential long-term storage stability when stored at lower temperatures.

2. Applicable Documents

2.1 ASTM Standards:

- D 270 Sampling Petroleum and Petroleum Products
- D 2274 Test for Oxidation Stability of Distillate Fuel Oil Accelerated Method
- D 1500 ASTM Color of Petroleum Products
- D 381 ASTM Jet Evaporation, Adherent Gums

3. Summary of Method

3.1 A measured volume of filtered fuel is stored in vented Pyrex bottles for three months at 110°F . After three months the samples are examined for color and sediment.

4. Significance

4.1 This test is the most commonly used accelerated test for distillate fuel oils. Many studies have been made and it has been shown that 13 weeks at 110°F dark storage is equivalent, in most cases, to one year of ambient storage (12.1).

5. Apparatus

5.1 Sample containers are Serum Pyrex Bottles (12.2). The total capacity of the container is 500 mL.

8. Preparation of Apparatus

- 8.1 Sample storage bottles. Rinse bottle with Adherent Insoluble solvent (equal parts of reagent grade toluene, acetone, methanol) and dry. If any stains are present, treat bottle first with chromic acid cleaning solution, rinse with water and dry.
- 8.2 Equilibrate the filter in a petri dish next to the balance for 30 minutes. Weigh filter to nearest 0.1 mg and retain for sample analysis. A "moisture blank" is run to correct for atmospheric moisture. Place the "moisture blank" filter in the petri dish, dry for 1 hour in an oven maintained at 194° ± 2°F. After drying place in desiccator (without desiccant) for at least 1 hour. One "moisture blank" filter is run for every four samples analyzed. The "moisture blank" filter should be weighed whenever the other four test filters are weighed and its change of weight subtracted from the level sample weight to correct for atmospheric moisture. The "moistur: blank" filter is equilibrated in a petri dish next to the balance for 30 minutes before weighing.

9. Preparation of Sample

9.1 Prior to storage, the fuel oil shall be filtered through Whatman No. 2 filter paper at room temperature. Determine ASTM D 1500 color on unaged filtered fuel.

10. Procedure

- 10.1 Place 350 mL of filtered fuel in a bottle. Label storage bottle. Place in Miracle Top Box (Sec. 5.2) and store in hot room at $110^{\circ} \pm 1^{\circ}$ F for 3 month dark storage.
- 10.2 At the end of the 3 month storage period remove the sample and cool to room temperature. Filter the aged sample under vacuum

(20-25 mm $\rm H_20$) through a weighed filter (0.8 μ Millipore Type AA) in a millipore filter holder. Remove the filtrate and determine ASTM D 1500 color on aged filtered fuel. Rinse the storage bottle with <u>n</u>-heptane to remove all traces of fuel from the bottle, transferring each wash to the filter holder. Wash the filter paper and holder with <u>n</u>-heptane until free of fuel oil, using vacuum. The rinse filtrate can be discarded.

10.3 Transfer the filter to a petri dish, dry for 1 hour in an oven maintained at 194° ± 2°F. After drying place in desiccator (without desiccant) for at least 1 hour. Equilibrate the filter together with the "moisture blank" next to balance for 30 minutes. Weight to nearest 0.1 mg.

ll. Calculations

11.1 Calculate the total sediment formation in milligrams per 100 mL after aging as follows:

$$A = \frac{B}{C} \times 100$$

where:

A = Total sediment, mg/100 mL

B = Weight of filterable sediment corrected for moisture using the "moisture blank"

C = Fuel placed in storage bottle, mL

12. References

12.1 Distillate Fuel Storage Stability Research conducted at Petroleum Experimental Station Bureau of Mines, Bartlesville, Oklahoma -- Done for Western Petroleum Refining Association, October, 1958.

- 12.2 Bottle, Storage, (Centrifuge, original form) Pyrex, Corning, New Jersey, No. 1260; CGW code 423290, 500 mL cap. 73 mm dia., 173 mm high.
- 12.3 Miracle Box No. 15, Paige Co., 432 Park Avenue South, New York, New York 10016.

A.4 Modification for Three Months 110°F Dark Storage Fuel Oil Stability Test

- Store samples in vented Pyrex bottle for three months at 110°F as per Three Months 110°F Dark Storage Stability Test.
- 2. After three months storage, place sample in sonic bath for 30 minutes.
- 3. Send samples for the following analysis:

C₇ Insolubles
Toluene Insolubles
Kinematic Viscosity at 100°F

- 4. Pour the remainder of sample into the bottle and store under N₂. Wash the sample containers with three rinses (of about 50 mL each) of cyclohexane. Discard cyclohexane rinse solution.
- 5. Dissolve any adherent gum on sample container wall with adherent insoluble solvent (equal parts of toluene, acetone and methanol). Determine the adherent insoluble content by evaporating the solvent at 320°F (160°C) by the air-jet method described in Method D 381. Obtain weight of adherent insolubles in mg to 0.1 mg.

A.5 Modification for F 21-61 (300°F Accelerated Fuel Oil Stability Test)

- 1. Follow steps (3), (4), and (5) of F 21-61 method.
- After sample has cooled (step 5), place in sonic bath for 30 minutes.
- 3. Send sample for the following analysis:

C₇ Insolubles
Toluene Insolubles
Kinematic Viscosity

4. Pour the remainder of sample into the bottle and store under N2.

A.6 Modification for ASTM D 2274-74

- 1. Follow step (8.3) of ASTM D 2274-74 method.
- After test, remove sample from bath and pour it into a 500 mL Erlenmeyer flask. Place the flask in a sonic bath for 30 minutes.
- 3. Sample for the following analysis:

C₇ Insolubles
Toluene Insolubles
Kinematic Viscosity at 100°F

- 4. Pour the remainder of sample into the bottle and store under N2.
- 5. Wash the sample container used in the ASTM D 2274-74 test and oxygen delivery tube with three rinsings (of about 50 mL each) of cyclohexane. Discard cyclohexane rinse solution.

6. Dissolve any adherent gum on container walls and oxygen delivery tube with the adherent insoluble solvent (equal parts of toluene, acetone and methanol). Determine the adherent insolubles content by evaporating the solvent at 320°F (160°C) by the air-jet method described in Method D 381. Obtain weight of adherent insolubles in mg to nearest 0.1 mg.

A.7 Standard Analytical Methods

	UOP	ASTM
Method Title	Method No.*	Method No.**
Distillation		D 1160
Heptane Insoluble Matter - Membrane Filter	U 614	
Toluene Insoluble Matter - Membrane Filter	Mod. U 614	
Conradson Carbon Residue of Petroleum Products		D 189
Viscosity, Kinematic at 100°F		D 445
Hydrogen and Carbon (Micro)	บ 638	
Oxygen, Total, in Organic Materials	U 649	
Sulfur in Petroleum Products		D 1512
Nitrogen by Acid Extraction & Kjeldahl (495)	U 384	
Steam Jet Gum		D 381
300°F Accelerated Fuel Oil Stability Test	Mod. DuPont	
	F 21-61	
Oxidation Stability of Distillate Fuel Oil		D 2274
Three Months 110°F Dark Storage Stability Test	***	
Metals in Petroleum Oil by AAS	800	
Bromine Number of Petroleum Distillates and		
Commerical Aliphatic Olefins by		
Electrometric Titration		D 1159
Modified Coulometric Sulfur	บ 731	

^{* &}quot;UOP Laboratory Test Methods for Petroleum and Its Products", UOP Process Division, Des Plaines, Illinois.

^{** &}quot;1980 Annual Book ASTM Standards", American Society for Testing and Materials, Philadelphia, PA; 1980.

^{***} UOP Laboratory Test Method, not yet finalized.

Appendix B.1

MAXIMUM JP-4 CASE -- PROFORMA FINANCIAL STATEMENT

ECONOMIC EVALUATION HODEL			USAF 984	SHALE OZL STUDY	ă		HAX.	MAX. JP4-15% ROI	Ħ	
RUN NJEGR 2		ĸ			STATEMENT				- PAGE	t t
ALL & VALUES IN MILLIONS	1985	1966	1961	1968	1969	1990	1661	1992	1993	7667
PERCENT CAPACITY	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
ANUAL SALES	665.76	1731.51	1731.51	1731.51	1731.51	1731.51	1731.51	1731.51	1731.51	1731.51
RAH MATERIAL COST	657.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00
GROSS MARGIN	208.76	417.51	417.51	417.51	417.51	417.51	417.51	417.51	417.51	417.51
OPERATING LABOR	8.80	9.80	8.60	9.80	8.80	6.80	9.90	9.00	9.80	6.60
UTILITY COST	10.19	20.38	20.38	20.38	20.38	20.38	20.39	20.38	80.38 20.38	20.38
CAT+CHEM COST	4.39	6.78 8.78	8.78 9.78	62.42 9.76	6.78 8.78	6.78 6.78	99 99	6.76	6.78	%.% •
DIRECT HEG EXPENSE	46.30	60.86	60.68	60.09	60.89	60.68	60.88	60.09	60.66	60.66
INGURANCE	4.41	4.61	6.41	4.41	4.41	4.41	4.61	4.41	4.61	4.41
PROPERTY TAX	8.63	6.63	9.83	9.83	9.63	6.63	9.63	9.03	0.03	9.93
INDIRECT MFG EXPENSE	13.24	13.24	13.24	13.24	13.24	13.24	13.24	13.24	13.24	13.24
DEPRECIATION INTEREST ON M. CAPITAL	112.00	103.38	13.8	66.15 13.95	7.52	68.92 13.95	60.31	51.69	13.98	. 3. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5.
TAXABLE INCOPE TAX LOSS DEDUCTION	23.26	226.05	234.67	243.28	251.90	260.51	269.13	277.74	286.36	294.97
ADJ. TAXABLE INCOME	23.26	226.05	234.67	243.28	251.90	260.51	260.13	277.74	286.36	24.97
CURRENT TAX TAX CREDITS	10.64	113.03	117.33	121.64	125.45 0.	130.26	134.56	139.67	143.18	147.49
TAXES PAYABLE	1.16	33.20	117.33	121.64	125.75	130.26	134.56	138.67	143.18	147.49
NET AFTER TAX INCOME	22.10	192.85	117.33	121.64	125.5	130.26	134.56	138.67	143.18	147.49
OPERATING CASH FLOW	134.10	2%.23	212.10	207.79	203.49	199.18	194.87	190.56	- 186,26 -	161.95
DEBT RETIREMENT	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
NET CASH FLOW	134.10	2%.23	212.10	207.79	203.49	199.18	194.87	190.56	186.24	181.95

ECONOMIC EVALUATION MODEL					i	
RUN NUTBER 2		ĸ	OFCORMA FIL	USAF SHALE UIL STUUT PROFORMA FINANCIAL STATEMENT	NEMENT	
ALL & VALUES IN MILLIONS	1995	1996	1997	1998	1999	2000
PERCENT CAPACITY	100.00	100.00	100.00	100.00	100.00	100.00
ANUAL SALES	1731.51	1731.51	1731.51	1731.51	1731.51	1731.51
RAH MATERIAL COST	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00
GROSS MARGIN	417.51	417.51	417.51	417.51	417.51	417.51
OPERATING LABOR	8.60	8.80	9.80	8.80	8.80	8.80
עזונוזי כספד	20.38	20.36	20.38	20.38	20.38	20.38
MAINTENANCE COST	22.92 A 78	22.92	22.92 av	22.92	22.92	22.92
	00	00	00	00	00	00
DIRECT HIG EXPENSE	60.88	60.98	60.88	60.68	60.86	60.99
INSURANCE	4.41	4.41	4.43	4.41	4.41	4.41
FROPERTY TAX	8.83	9.93	0.63	6.63	9.93	8.83
INDIRECT HFG EXPENSE	13.24	13.24	13.24	13.24	13.24	13.24
DEPRECIATION	25.85	17.23	8.62			
INTEREST ON M. CAPITAL	11.95	13.95	13.95	13.95	13.8	13.95
TAXABLE INCOME TAX LOSS DEDUCTION	303.59	312.20	320.02	329.44	329.44	329.44
ADJ. TAXABLE INCOME	303.59	312.20	320.62	329.44	329.44	329.44
CURPENT TAX TAX CREDITS	151.79 0.	156.10 0.	160.41	164.72	164.72 0.	164.72 0.
TAXES PAYABLE	151.79	156.10	160.41	164.72	164.72	164.72
NET AFTER TAX INCOME	151.79	156.10	160.41	164.72	164.72	164.72
OPERATING CASH FLOW	177.64	173.33	169.03	164.72	164.72	164.72
DEBT RETIREMENT RETURN OF HORK. CAP.	.	ó	ė	ó	ó	0. 116.00
NET CASH FLOH	177.64	173.33	169.03	164.72	164.72	164.72

of the second of

MAX. JP4-15% ROI

PROFITABILITY SUPPARY

ECONOMIC EVALUATION MODEL RUN NUMBER 2

ALL & VALUES IN MILLIONS

PAYBACK PERIOD: YEARS

4.26

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON EQUITY X

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON PROJECT %

101.09

NET PRESENT VALUE 315%: 6HH

0.11

751e

N.P.V. INDEX

USAF SPALE OIL STUDY

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PROFITABILITY SUPPLARY

BASIS FOR STUDY

	10000000000000000000000000000000000000	\$6.82 \$6.82 155.00	93. 0. 0. 0. 0. 0. 0.
ECONOMIC EVALUATION MODEL RUN NAMBER 2 ALL \$ VALUES IN MILLIONS FORMANT PABAMETERS		BASE FEED PRICES SHALE OIL BASE PRODUCT PRICES JET FUEL 4/BBL APPONIA 4/ST SULFUR 4/LT	BASE HORKING CAPITAL RAW MATERIALS ACCUONTS PAYABLE ACCOUNTS PAYABLE NOBLE HETALS LAND CATALYSTACHEHICALS ROYALTY & FEES

FOUNDATE EVALUATION MODEL			USAF SH	USAF SHALE OIL STUDY			
RUN NUBER 2			INVESTM	INVESTMENT SUTHARY	MAX. JP4-15% R01	PAGE	4
ALL * VALUES IN MILLIONS							
ESTIMATED CAPITAL INVESTMT CONST. INFLATION FACTOR INTEREST RATE: %	784.00 1.000 0.000						
YEAR	1962	1961	1984				
% COMPLETION FOR YR	25.00	50.00	25.00				
PLANT INVESTMENT . CAPITAL INVESTMENT	196.00	392.00	196.00			1	
. CONST. COST INFLATION	-0.00	-0.00	-0.00				
. CONST. COST FOR YEAR	196.00	392.00	196.00				
TOTAL DEPRECIABLE INV.				784.00			
OTHER INVESTMENT LAND INITIAL CAT.£CHEM. ROYALTY & FEES				25.00 0.00		•	
SUBTOTAL				25.00			
HORKING CAPITAL RAH HATERIALS ACCOUNTS RECEIVABLE ACCOUNTS PAYBLE NOSLE METALS INT ON CONSTRUCTION	ó	ó	ó	93.00 0.0 0.0	· :	!	
TOTAL CONST INTEREST SUBTOTAL				93.00			
TOTAL INVESTMENT				902.00			

USAF SHALE OIL STUDY

Appendix	B.1	(Cont	.)
Appendia	D •••	(00	• /

	ECONOMIC EVALUATION MODEL					ì		HAX.	MX. JP4-15% ROI	20	
	RUN NUMBER 2 ALL & VALUES IN MILLIONS			USAF SHALE OF PRODUCTION	ile oil study Or: Su pr iary	.				PAGE	5 - 1
	YEAR	1985	1986	1967	1988	1989	1990	1991	1992	1993	1994
		50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
	PRODUCTION, IT UNITS/YEAR JET FUEL (BBLS)	15.07	30.14	30.14	30.14	30.14	30.14	30.14	30.14	30.14	30.14
-	SULFUR (LT)	0.02	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.03	0.10
	PRODUCT PRICES: #/UNIT JET FUEL \$/68L	56.82	56.82	56.82	56.82	56.82	56.82	56.82	56 .82	26.95	56.82
	ATTONIA 8/ST SULFUR 8/LT	155.00 105.00	155.00	155.00	155.00	155.00 105.00	155.00	155.00	155.00	155.00 105.00	155.00 105.00
	FRODUCT VALUES OHY/YEAR JET FUEL ATTORIA SULFUR	856.42 2.86 0.83	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66

ECONOMIC EVALUATION MODEL							MAX. JP4-15% ROI		
RUN NUMBER 2 ALL & VALUES IN MILLIONS			USAF SHALE O	ale oil study Ion suppary	<u>`</u>			PAGE	ru 1
YEAR	1995	1996	1997	1998	1999	2000			
	100.00	100.00	100.00	100.00	100.00	100.00			
PRODUCTION, PHI UNITS/YEAR JET FUEL (BBLS) ATTONIA (ST) SULFUR (LT)	30.14 0.10 0.03	30.14 0.10 0.03	30.14 0.10 0.03	30.14 0.10 0.03	30.14 0.10 0.03	30.14 0.10 0.03	;		
PRODUCT PRICES: \$/UNIT JET FUEL \$/BBL APPONIA \$/ST SULFUR \$/LT	56.82 155.00 105.00	56.82 155.00 105.00	56.82 155.00 105.00	56.82 155.00 105.00	56.82 155.00 105.00	56.82 155.00 105.00	:		•
PRODUCT VALUES BHLYTEAR JET FUEL AMTOKLA SULFUR	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.85 5.72 1.66	1712.65 5.72 1.66			

Appendix B.2

JP-4 PLUS DIESEL CASE -- PROFORMA FINANCIAL STATEMENT ECONOMIC EVALUATION MODEL

ECONOMIC EVALUATION HODEL	Jr-4 ruus 1	DIESEL CASE	! !	€	r ingavetal	, otheren	_	JP4+DIESEL-15% ROI	ROI	
RUN NUMBER 2		ă.	USAF SHA PROFORMA FIN	SHALE OIL STU FINANCIAL STA	STUDY				PAGE	1 - 1
ALL & VALUES IN MILLIONS	1965	1986	1961	1988	1989	1990	1661	1992	1661	1994
PERCENT CAPACITY	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
AMUAL SALES	886.76	1773.53	1773.53	1773.53	1772.53	1773.53	1773.53	1773.53	1773.53	1773.53
RAW MATERIAL COST	657.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00
GROSS MARGIN	229.76	459.53	459.53	459.53	459.53	459.53	459.53	459.53	459.53	459.53
OPERATING LABOR	96.6	96.6	90.0	96.08	96.6	6	0	0	80.0	0
UTILITY COST	13.29	26.58	26.58	26.58	26.58	26.58	26.58	26.58	26.58	26.58
MAINTENANCE COST	29.63	25.62	25.62	25.62	25.62	25.62	25.62	25.62	25.62	25.62
CAT+CHEM COST	4.06	8.11	8.11	8.11	8.11	8.11	6.11	6.11	8.11	9.11
DIRECT MFG EXPENSE	52.94	70.29	70.29	70.29	70.29	70.29	70.29	70.29	70.29	70.29
INSURANCE	4.85	4.85	4.85	4.85	4.85	4.85	4.85	4.85	4.85	4. A5
PROPERTY TAX	9.70	9.70	9.70	9.70	9.70	9.70	9.70	9.70	9.70	9.70
INDIRECT MFG EXPENSE	14.53	14.55	14.55	14.55	14.55	14.55	14.55	14.55	14.55	14.55
DEPRECIATION	124.66	115.25	105.65	8.04	96.44	76.84	67.23	57.63	48.02	38.42
INTEREST ON M. CAPITAL	13.05	13.95	13.95	13.95	13.95	13.95	13.95	13.95	13.95	13.95
TAXABLE INCOME TAX LOSS DEDUCTION	23.46	245.49	255.10	264.70	274.30	283.91	293.51	303.12	312.72	322.33
ADJ. TAXABLE INCOME	23.46	245.49	255.10	264.70	274.30	263.91	293.51	303.12	312.72	322.33
CURPENT TAX TAX CREDITS	11.73	122.75 88.43	127.55	132.35	137.15	141.95	146.76	151.56 0.	156.36	161.16
TAXES PAYABLE	1.17	34.32	127.55	132.35	137.15	141.95	146.76	151.56	156.36	161.16
NET AFTER TAX INCOME	22.29	211.18	127.55	132.35	137.15	141.95	146.76	151.56	156.36	161.16
OPERATING CASH FLOW	147.15	326.43	233.20	228.39	223,59	218.79	213,99	209, 18	204.38	199.58
OEBT RETIREHENT	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
NET CASH FLOW	147.15	326.43	233.19	228.39	223.59	218.79	213.99	209.18	204.38	199.58

	ECONOMIC EVALUATION MODEL							JP4+D]
	RUN NUMBER 2		δ.	USAF SHI ROFORMA FI	USAF SHALE OIL STUDY PROFORMA FINANCIAL STATEMENT	JOY NTEMENT		
	ALL & VALUES IN MILLIONS YEAR	1995	19%	1997	1998	1999	2000	
	PERCENT CAPACITY	100.00	100.00	100.00	100.00	100.00	100.00	
	ANUAL SALES	1773.53	1773.53	1773.53	1773.53	1773.53	1773.53	
	RAH MATERIAL COST	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	
	GROSS MARGIN	459.53	459.53	459.53	459.53	459.53	459.53	
	OPERATING LABOR	96.6	9.08	9.98	9.98	0	9.98	
	UTILITY COST	26.58	26.58	26.58	26.58	26.58	26.58	
	MAINTENANCE COST CAT+CHEM COST	25.62	25.62	25.62	25.62	25.62	25.62	
		7.0	44.0	11.0	77.0	11.0	11:0	
•	DIRECT HIG EXPENSE	70.29	70.29	70.29	70.29	70.29	70.29	
-18	INSURANCE	4.85	4.85	4.85	4.85	4.85	4.85	
9-	PROPERTY TAX	9.70	9.70	9.70	9.70	9.70	9.70	
	INDIRECT HFG EXPENSE	14.55	14.55	14.55	14,55	14.55	14.55	
	DEPRECIATION INTEREST ON M. CAPITAL	28.81 13.95	19.21	9.61	13.95	13.95	13.95	
	TAXABLE INCOME TAX LOSS DEDUCTION	331.93	341.53	351.14	360.74	360.74	360.74	
	ADJ. TAXABLE INCOME	331.93	341.53	351.14	360.74	360.74	360.74	
	CURRENT TAX TAX CREDITS	165.97	170.77 0.	175.57	180.37	180.37	180.37 0.	
	TAXES PAYABLE	165.97	170.77	175.57	180.37	180.37	180.37	
	NET AFTER TAX INCOME	165.%	170.77	175.57	160.37	160.37	180.37	
	OPERATING CASH FLOW	194.78	189.98	2t'50t	180.37	160.37	180.37	
	DEBT RETIREMENT PETURN OF WORK. CAP.	ó	ó	ó	ó	ö	0. 115.00	
	NET CASH FLOW	194.78	169.98	185.17	160.37	160.37	180.37	

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PROFITABILITY SUPPIARY

ALL & VALUES IN MILLIONS

PAYBACK PERIOD: YEARS

15.00 AFTER TAX DISCOUNTED RATE OF RETURN OF PROJECT X AFTER TAX DISCOUNTED RATE OF RETURN ON EQUITY 2

111.05 NET PRESENT VALUE 215%: 91%

N.P.V. INDEX

915%

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ECONOMIC EVALUATION MODEL RUN NJ. BER 2

	50.000 0.000 0.000 1.000 1.000 1.000	10.00 90.000	60.00	58.82 155.00 105.00	93.00
ALL . JALUES IN MILLIONS ECONOTIC PARAMETERS	ELT LIFE: YEAR TREST RATE: X "E TAX RATE: ENT DEBT ALL INFLATION TY INPLATION TY INPLATION TY INPLATION TY OBOST INFL TOOST INFL THE COST INFL	INV. TAX CREDIT: X OF CAP INV TX CR: MAX X OF TAX 61 1 BASE FEED FRICES	SHALE OIL RASE PRODUCT PRICES	JET FUEL \$/EBL APPONIA \$/ST SULFUR \$/LT BASE HORKING CAPITAL	RAW MATERIALS ACCUDITS RECEIVABLE ACCOUNTS PAYABLE NOBLE METALS LAND CATALYSTECHEMICALS **OYALTY & FEES

ECONOMIC EVALUATION MODEL RUN NUMBER 2			INVESTHEN	INVESTHENT SUPPARY	JP4+DIESEL-15% ROI	PAGE	t - 5
ALL & VALUES IN MILLIONS							
ESTIMATED CAPITAL INVESTMT CONST. INFLATION FACTOR INTEREST RATE: X	874.00 1.000 0.000						
YEAR	1982	1981	1984				!
2 COMPLETION FOR YR	25.00	50.00	25.00				
PLANT INVESTHENT . CAPITAL INVESTHENT	218.50	437.00	218.50			•	 -
. CONST. COST INFLATION	-0.00	-0.00	-0.00				
. CONST. COST FOR YEAR	218.50	437.00	218.50				
TOTAL DEPRECIABLE INV.				874.00			
OTHER INVESTHENT LAND . INITIAL CAT. ECHEM ROYALTY & FEES				22.00 0.00	!	1	
SUBTOTAL				22.00			
HORKING CAPITAL RAH HATERIALS ACCOUNTS RECEIVABLE ACCOUNTS PAYABLE				00.00 0.00	1		
. NOSLE PETALS . INT ON CONSTRUCTION TOTAL CONST INTEREST	ó	ó	ó	<i>.</i>			i
SUBTOTAL				93.00			;
TOTAL INVESTMENT				00.999			

			LISAF SHA	ILE OTI STU	č		****	JP4+U1E3EL-157 KU1	108		
RUN NUMBER 2 ALL & VALUES IN MILLIONS			PRODUCTION	ON SUPPLARY	5				PAG	PAGE 5 - 1	
YEAR	1985	1986	1967	1986	1989	0661	1991	1992	1993	1994	
	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	
PRODUCTION, MH UNITS/YEAR JET FUEL (BBLS)	14.92	29.84	29.84	29.84	29.84	29.84	29.84	29.84	29.84	29.84	
APPONIA (ST.)	0.05	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	
	0.01	0.03	0.03	0.03	0.03	0.03	0.03	0.03	0.0 0.0	0.03	
PRODUCT PRICES: \$/UNIT JET FUEL \$/88L	58.82	58.82	59.82	58.82	58.82	59.62	58.82	58.82	58.82	88.	
AMMONIA \$/ST	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	
SULFUR 6/LT	105.00	105.00	105.00	105.00	105.00	105.00	105.00	105.00	105.00	105.00	
PRODUCT VALUES SHYYEAR		ì	ì	į		i	į	į	;		
APTOLIA	8.7. 8.7.	26.5 5.92	5. 35	1/54.86 5.92	1754.86 5.92	1754.86 5.92	1754.86 5.92	1754.86 5.92	1754.86	1,754.86	- 1
SULFUR	98.0	1.71	1.71	1.71	1.71	1.71	1.71	1.71	1.71	1.71	

ECONOMIC EVALUATION MODEL			AND SHA	IE OTI STU	`		JP4+DIESEL-15% ROI
RUN NUMBER 2 ALL \$ VALUES IN MILLIONS			PRODUCTI	PRODUCTION SUPPARY	, 5		
YEAR	1995	10%	1997	1998	1999	2000	
	100.00	100.00	100.00	100.00	100.00	100.00	
PRODUCTION, HT UNITS/YEAR JET FUEL (BBLS) APPONIA (ST) SULFUR (LT)	29.84 0.10 0.03	29.84 0.10 0.03	29.84 0.10 0.03	29.84 0.10 0.03	29.84 0.10 0.03	29.84 0.10 0.03	:
PRODUCT PRICES: \$/UNIT JET FUEL \$/BBL AFFORMA \$/ST SULFUR \$/LT	58.82 155.00 105.00	58.82 155.00 105.00	58.82 155.00 105.00	58.82 155.00 105.00	59.62 155.00 105.00	58.82 155.00 105.00	
PRODUCT VALUES \$P\$VYEAR JET FUEL APPONIA SULFUR	1754.86 5.92 1.71	1754.86 5.92 1.1	1754.86 5,92 1.1	1754.86 5.92 1.71	1754.86 5.92 1.71	1754.86 5.92 1.71	

Appendix B.3

MAXIMUM JP-8 CASE -- PROFORMA FINANCIAL STATEMENT

ECONOMIC EVALUATION HODEL					į		HAX	HAX. JP8-15% ROI	10	
RUN NUBER 2		K	USAF SHALE OIL STUDI PROFORMA FINANCIAL STATEMENT	SHALE OIL STOOT	JOY NTEPENT		,		PAGE	1 - 1
ALL & VALUES IN MILLIONS YEAR	1965	1986	1987	1988	1989	1990	1661	1992	1993	1994
PERCENT CAPACITY	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
ANNUAL SALES	874.55	1749.10	1749.10	1749.10	1749.10	1749.10	1749.10	1749.10	1749.10	1749.10
RAH HATERIAL COST	657.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00	1314.00
GROSS MARGIN	217.55	435.10	435.10	435.10	435.10	435.10	435.10	435.10	435.10	435.10
OPERATING LABOR	10.16	10.16	10.16	10,16	10.16	10.16	10.16	10.16	10.16	10.16
UTILITY COST	10.68	21.36	21.36	21.36	21.36	21.36	21.36	21.36	21.36	21.36
CATECHEM COST	23.97	23.97 8.98	23.97 6.98	23.97 8.98	23.97 8.98	23.97 8.98	23.97	23.97	6.88	23.97
DIRECT MFG EXPENSE	49.30	64.47	64.47	64.47	64.47	64.47	64.47	64.47	64.47	64.47
INSURANCE PROPERTY TAX	4.60	4.60 9.20	4.60 9.20	4.60	4.60 9.20	4.60 9.20	4.60 9.20	4.60	4.60	4.60
INDIRECT HEG EXPENSE	13.60	13.80	13.60	13.60	13.60	13.60	13.60	13.80	13.60	13.80
DEPRECIATION INTEREST ON H. CAPITAL	117.00	108.00 13.95	99.00	90.00	61.00 13.95	72.00	63.00	54.00	45.00	36.00
TAXABLE INCOME TAX LOSS DEDUCTION	23.50	234.88	243.68	252.88	261.88	270.88	279.68	288.88	297.86	306.66
ADJ. TAXABLE INCOME	23.50	234.88	243.80	252.88	261.88	270.88	279.86	288.00	297.68	306.88
CURRENT TAX TAX CREDITS	11.75	117.44	121.94 0.	126.44 0.	130.94 0.	135.44	139.94	144.44	146.94	153.44
TAXES PAYABLE	1.17	34.02	121.94	126.44	130.96	135.44	139.96	166.46	146.94	153.44
NET AFTER TAX INCOME	22.33	200.65	121.94	126.44	130.9%	135.44	139.94	164.44	140.94	153.44
OPERATING CASH FLOW	159.33	108.85	220.94	216.44	211.94	207.44	202.94	198.44	193.94	189.44
DEBT RETIRENENT	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
NET CASH FLOW	139.33	306.85	220.94	216.44	211.94	207.44	202.94	199.44	193.94	189.44

Appendix B.3	(Cont	•)
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ECONDMIC EVALUATION MODEL							MAX. JP6-15% ROI
RUN NUMBER 2		æ	USAF SHALE DIL STUDY PROFORMA FINANCIAL STATEMENT	USAF SHALE DIL STUDY ORMA FINANCIAL STATE	JOY ATEMENT		
ALL & VALUES IN MILLIONS YEAR	1995	19%	1997	1998	1999	2000	
PERCENT CAPACITY	10,.00	100.00	100.00	100.00	100.001	100.00	
ANNIAL SALES	1747.10	1749.10	1749.10	1749.10	1749.13	1749.13	
RAH HATERIAL COST	1314.00	1,514.00	1314.00	1314.00	1314.30	1314 20	
GROSS MARGIN	435.10	435.10	435.10	+35.10	435.10	435.10	
OPERATING LABOR	4. 9.	71 91	10.00	**		;	
UTILITY COST	21.36	21.36	21.36	21.36	20.16	10. 16 21 14	
MAINTENANCE COST	23.97	23.97	23.97	23.97	23.97	23.97	
CAT+CHEM COST	9 . 30	9 .8	9	8.48	8.8	6.98	
DIRECT MFG EXPENSE	64.47	64.47	64.47	64.47	64.47	64.47	
INSURANCE	4.60	4.60	4,60	4.60	9	04.4	
PROPERTY TAX	9.20	9.50	9.50	9.20	9.20	9.20	
INDIRECT MFG EXPENSE	13.80	13.80	13.60	13.80	13.60	13.80	
DEPRECIATION INTEREST ON M. CAPITAL	27.00	18.00	9.00	13,95	13.95	13, 95	
TAXABLE INCOME TAX LOSS DEDUCTION	315.88	324.88	333.68	342.88	342.88	342.88	
ADJ. TAXABLE INCOME	315.88	324.68	333.68	342.88	342.88	342.88	
CURRENT TAX TAX CREDITS	157.94 0.	162.44	166.94 0.	171.44	171.44	171.44	
TAXES PAYABLE	157.94	162.44	166.94	171.44	171.44	171.44	
NET AFTER TAX INCOME	157.94	162.44	166.94	171.44	171.44	171.44	
OPERATING CASH FLOW	184.94	180.44	175.94	171.44	171.44	171.44	1
OEBT RETIREMENT RETURN OF NORK. CAP.	ö	ö	ö	ó	ö	0. 120.00	
NET CASH FLOH	184.94	180.44	175.94	171.44	171.44	171.44	

MAX. JP8-15% ROI

USAF SHALE OIL STUDY

PROFITABILITY SUPPARY

ECONOMIC EVALUATION MODEL RUN NUMBER

ALL & VALUES IN MILLIONS

PAYBACK PERIOD: YEARS

4. 26

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON EQUITY X

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON PROJECT 2

105.25

NET PRESENT VALUE 315%: 8159

0.11

251e

N.P.V. INDEX

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15%	

PROFITABILITY SUPPLARY

BASIS FOR STUDY

ECUMPIC EVALUATION MODEL RUN NUMBER 2	
ALL # VALUES IN MILLIONS	
ECONOMIC PARAMETERS	
	16
LUNSIM PERIOD: YEARS INTEREST RATE: %	0.000 •
INCOME TAX RATE: X PERCENT DEBT	50.000
GENERAL INFLATION FACTOR	1.000
LABOR INFLATION FACTOR	1.000
UTILITY INFLATION FACTOR	000.
CORIST COST INFL FACTOR	000.1
-	1.000
INV. TAX CREDIT: % OF CAP	10.00
INV TX CR: MAX % OF TAX	90.00
BASE FEED PRICES	
SHALE OIL	40.00
BASE PRODUCT PRICES	
JET FUEL \$788L APPONIA \$7ST	58.80 155.00
SULFUR 6/LT	205 00

RAW HATERIALS
ACCUONTS RECEIVABLE
ACCOUNTS PAYABLE
NOBLE HETALS
LAND
CATALYSTECHMICALS
ROYALTY & FEES

93.00 0. 0. 27.00

-198-

919.00

27.00 0. 27.00

LAND INITIAL CAT. SCHEM. ROYALTY & FEES

SUBTOTAL

OTHER INVESTMENT

93.00

ö

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HORKING CAPITAL
RAH HATERIALS
ACCOUNTS RECEIVABLE
ACCOUNTS PAYABLE
NOBLE METALS
INT ON CONSTRUCTION
TOTAL CONST INTEREST

TOTAL INVESTMENT

SUBTOTAL

ECONOMIC EVALUATION MODEL			USAF SHA	USAF SHALE OIL STUDY		
RUN PAPBER 2			INVESTME	INVESTMENT SUPPARY	MAX. JP8-15% ROI	<u>u</u>
ALL & VALUES IN MILLIONS						Ž.
ESTIMATED CAPITAL INVESTMI CONST. INFLATION FACTOR INTEREST RATE: X	81 3. 00 1 300 0. 000					į
YEAR	1982	1961	1984			
% COMPLETION FOR YR	25.00	50.00	25.00			
PLAIT INVESTHENT CAPITAL INVESTHENT	204.75	409.50	204.75			
CONST. COST INFLATION	-0.00	-0.00	-0.00		. !	· · · · · · · · · · · · · · · · · · ·
TOTAL DEPRECIABLE INV.	504.75	409.50	204.75	919.00		

RLN NLIBER 2 ALL 4 VALUES IN MILLIONS			USAF SHALE OF SPRODUCTION S	NLE OIL STUDY ION SUPPARY	à .		T.	MAX. JPG-15% ROI	T PAGE	•	;
YEAR	1965	1986	1987	1988	1989	1990	1661	1992	1991	1994	:
	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	
PRODUCTION, HI UNITS/YEAR JET FUEL (BBLS)	14.72	29.43	29.43	29.43	29.43	29.43	29.43	29.43	29.43	29.43	İ
SULFUR (LT)	0.0	0.03	0.03	0.03	0.03	0.00	0.03	0.10	0.0	0.03	ļ
	58.60	58.80	58.80	56.60	56.60	58.80	56.60	58.60	56.60	56.60	į
SULFUR 6/LT	155.00 105.00	155.00 105.00	155.00 105.00	155.00	155.00 105.00	155.00 105.00	155.00 105.00	155.00 105.00	155.00 105.00	155.00 105.00	
PRODUCT VALLES SPYYEAR JET FUEL APPORTA	865.19 2.97	1730.38 5.94	1730.38 5.94	1730.38 5.94	1730.56 5.94	1730.38 5.94	1730.38 5.94	1730.38	1730.36	1730.38	:
5	8	**	**	7:1	7: 7	*:	7	1.1	7:1	7	

ECONOMIC EVALUATION MODEL							MAX19815% BOT	
RUN NUMBER 2 ALL & VALUES IN MILLIONS			USAF SHALE (PRODUCTION !	ALE OIL STUDY ION SUPPARY	Ď.			PAGE
YEAR	1995	19%	1997	1998	1999	2000		
	100.00	100.00	100.00	100.00	100.00	100.00		
PRODUCTION, MY UNITS/YEAR JET FUEL (BBLS) AMTONIA (ST) SULFUR (LT)	29.43 0.10 0.03	29.43 0.10 0.03	29.43 0.10 0.03	29.43 0.10 0.03	29.43 0.10 0.03	29.43 0.10 0.03		
PRODUCT PRICES: #/UNIT JET FUEL #/BBL AFFORIA #/ST SULFUR #/LT	58.80 155.00 105.00	58.80 155.00 105.00	58.60 155.00 105.00	56.60 155.00 105.00	\$6.80 155.00 105.00	58.60 155.00 105.00		
PRODUCT VALUES BITLYEAR JET FUEL ANTONIA SULFUR	1730.38 5.94 1.71	1730.38 5.94 1.71	1730.38 5.94 1.71	1730.38 5.94 1.71	1730.38 5.94 1.71	1730.38 5.94 1.71	:	

Appendix B.4

JP-8 PLUS DIESEL CASE -- PROFORMA FINANCIAL STATEMENT

ECONOMIC EVALUATION MODEL							JP8•	JP8+DIESEL-15% ROI	ROI	
RUN NUMBER 2		Æ	USAF SHALE OIL PROFORMA FINANCIAL		STATEMENT				PAGE	1 - 1
ALL & VALUES IN MILLIONS YEAR	1985	1986	1961	1988	1989	1990	1991	1992	1993	7661
PERCENT CAPACITY	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
ANUAL SALES	865.94	1731.68	1731.68	1731.68	1731.68	1731.68	1731.88	1731.88	1731.68	1731.68
RAH MATERIAL COST	657.00	1314.00	1314.00	1314.00	1314.00	1314.00	0	1314.00	1314.00	1314.00
GROSS MARGIN	208.94	417.68	417.88	417.88	417.88	417.88	417.88	417.88	417.88	417.88
OPERATING LABOR	10.16	10.16	10.16	10.16	10.16	10.16	10.16	10.16	10.16	10.16
UTILITY COST	9.32	18.64	18.64	18.64	18.64	18.64	18.64	18.64	18.64	18.64
MAINTENANCE COST CAT+CHEM COST	23.10	23.10 A 17	23.10 A 17	23.10 8.17	23.10 A 17	23.10	23.10 A 17	23.10 A.17	23.10 A 17	23.10 8.17
		78.0	77.0	76.0	16.0	15.0	76.0	75.0	15.0	100
DIRECT MFG EXPENSE	46.77	60.27	60.27	60.27	60.27	60.27	60.27	60.27	60.27	60.27
INSURANCE	4.43	4.43	4.43	4.43	4.43	4.43	4.43	4.43	4.43	4.43
PROPERTY TAX	8.86	8.86	8.8	8.86	8.86	8.8	8.8	8.8	8. 8	9.86
INDIRECT MFG EXPENSE	13.29	13.29	13.29	13.29	13.29	13.29	13.29	13.29	13.29	13.29
DEPRECIATION	112.86	104.18	95.49	86.81	78.13	69.45	60.77	52.09	43.41	34.73
INTEREST ON W. CAPITAL	13.80	13.60	13.80	13.80	13.80	13.80	13.80	13.80	13.80	13.60
TAXABLE INCOME TAX LOSS DEDUCTION	22.23	226.34	235.02	243.70	252.38	261.06	269.75	278.43	287.11	295.79
ADJ. TAXABLE INCOME	22.23	226.34	235.02	243.70	252.38	261.06	269.75	278.43	267.11	295.79
CURRENT TAX TAX CREDITS	11.11	113.17	117.51	121.85	126.19	130.53	134.67	139.21	143.55	147.89
TAXES PAYABLE	1.11	32.58	117.51	121.85	126.19	130.53	134.87	139.21	143.55	147.89
NET AFTER TAX INCOME	21.12	193.76	117.51	121.85	126.19	130.53	134.87	139.21	143.55	147.89
OPERATING CASH FLOH	133.97	297.93	213.00	208.66	204.32	199.98	195.64	191.30	386.98	182.62
DEBT RETIREMENT	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
NET CASH FLOW	133.97	297.93	213.00	208.66	204.32	199.98	195.64	191.30	186.%	182.62

A 1997年,1997年,1997年,1998年,199

JP8+DIESEL-15% ROI	0.00	100.00	1731.68	1314.00	417.88	10.16	16.64	23.10 3.37	69.27	87.7	8.86	13.29	00	00.01	330.51	330.51	165.26 0.	165.26	165.26	165.26	0. 115.00	165.26
	o			_	;				i	m	•			;				1			ä	;
JOY	1999	100.00	1731.88	1314.00	417.89	10.16	18.64	23.10	60.27	4.43	8 .86	13.29	- C	0.,1	330.51	330.51	165.26	165.26	165.26	165.26	ö	165.26
SHALE OIL STUDY FINANCIAL STATE	1998	100.00	1731.88	1314.00	417.88	10.16	18.64	23.10 8.37	60.27	4.43	8.	13.29	- A	20:51	330.51	330.51	165.26 0.	165.26	165.26	165.26	ö	165.26
USAF SHALE OIL STUDY PROFORMA FINANCIAL STATEMENT	1997	100.00	1731.88	1314.00	417.89	10.16	18.64	23.10 6.37	60.27	4.43	8.	13.29	69 3	20.51	321.83	321.03	160.92	160.92	160.92	169.60	ó	169.60
ã.	19%	100.00	1731.88	0	417.88	10.16	18.64	23.10 8.37	60.27	4.43	9.86	13.29	17.36		313 15	315.13	156.58 0.	156.58	156.58	173.94	ó	173.94
	1995	100.00	1731.88	1314.00	417.88	10.16	18.64	23.10 8.37	60.27	4.43	8)	26.04		304.47	304.47	152.24	152.24	152.24	178.28	ö	178.28
ECONOMIC EVALUATION MODEL RUN NUMBER 2	ALL & VALUES IN MILLIONS	PERCENT CAPACITY	antual sales	RAH MATERIAL COST	GROSS MARGIN	OPERATING LABOR	UTILITY COST	CAT+CHEM COST	DIRECT MFG EXPENSE		TACPER I PAX	INDIRECT MFG EXPENSE	DEPRECIATION INTEREST ON W. CAPITAL		TAXABLE INCOME TAX LOSS DEDUCTION	ADJ. TAXABLE INCOME	CURPENT TAX TAX CREDITS	TAXES PAYABLE	NET AFTER TAX INCOME	OPERATING CASH FLOH	OEBT RETIREMENT RETURN OF HORK. CAP.	HET CASH FLOH
										-20	3-											

The state of the s

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JP8+DIESEL-15% ROI

PROFITABILITY SUPPARY

USAF SHALE OIL STUDY

ECONOMIC EVALUATION MODEL RUN NUMBER 2

ALL & VALUES IN MILLIONS

PAYBACK PERIOD: YEARS

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON EQUITY X

4.26

101.44

NET PRESENT VALUE 215%: 617

0.11

215%

N.P.V. INDEX

15.00

AFTER TAX DISCOUNTED RATE OF RETURN ON PROJECT 2

-204-

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STG	
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BASIS	
a	

PROFITABILITY SUPPARY

JP8+DIESEL-15% ROI

			16	00.00	50.000	0.00	1.000	1.000	1.000	1.000	1.000	1.000	10.00	90.000
ECCHONIC EVALUATION MODEL RUN NUMBER 2	ALL & VALUES IN MILLIONS	ECONOMIC PARAMETERS	PROJECT LIFE:YEARS	INTEREST RATE: %	INCOME TAX RATE: 2	FERCENT DEBT	GENERAL INFLATION FACTOR	LABOR INFLATION FACTOR	UTILITY INFLATION FACTOR	PETRO PROOS INFL FACTOR	CONST COST INFL FACTOR	HORKING CAP INFL FACTOR	INV. TAX CREDIT: X OF CAP	INV TX CR: HAX X OF TAX
														-2

	40.00
PRICES	
BASE FEED	SHALE OIL

	RICES	
SHALE OIL	BASE PRODUCT PRICES	JET FUEL \$789L

58.85	155.00 105.00	
JET FUEL \$/BBL	ANTONIA \$/ST SULFUR \$/LT	

BASE WORKING CAPITAL	HATERIALS 92.00	ACCUONTS RECEIVABLE 0.	PAYABLE	LE METALS 0.		CATALYST&CHEMICALS 23.00	ALTY & FEES 0.
BASE MORKIN	RAH MATERIALS	ACCUONTS RE	ACCOUNTS PV	NOBLE METALS	LAND	CATALYST&CH	ROYALTY & F

	<u>"</u>				ທ	
RAM MATERIALS	ACCUONTS RECEIVABLE	ACCOUNTS PAYABLE	NOBLE METALS	LAND	CATALYSTACHEMICALS	DOVALTY & CCCC

ECCHONIC EVALUATION MODEL RUN NUMBER 2			USAF SHALE OIL STUINESTUINESTHENT SUFFFARY	USAF SHALE OIL STUDY INVESTHENT SUFFIARY	JP8+DIESEL-15% ROI	PAGE	4 - 1	
ALL \$ VALUES IN MILLIONS								
ESTIMATED CAPITAL INVESTMT CONST. INFLATION FACTOR INTEREST RATE: %	790.00 1.000 0.000							
YEAR	1982	1983	1984				!	
% COMPLETION FOR YR	25.00	50.00	25.00					
PLANT INVESTHENT . CAPITAL INVESTHENT	197.50	395.00	197,50					
. CONST. COST INFLATION	-0.00	-0.00	-0.00					
. CONST. COST FOR YEAR	197.50	395.00	197.50					
TOTAL DEPRECIABLE INV.				790.00				
OTHER INVESTMENT . LAND . INITIAL CAT. & CHEM ROYALTY & FEES				23.00 0.			\$	1
SUBTOTAL			•	23.00			:	
HORKING CAPITAL . RAH MATERIALS . ACCOUNTS RECEIVABLE . ACCOUNTS PAYABLE . NOBLE METALS . INT ON CONSTRUCTION TOTAL CONST INTEREST	ò		ó	92.00 0.0 0.0				
SUBTOTAL				05.00		,	;	
TOTAL INVESTMENT				905.00				

LIGHT EVALUATION TO THE							à	JP8+DIESEL-157 KO	5		
RUN NUMBER 2 ALL & VALUES IN MILLIONS			USAF SHALE O	le oil study On su ri ary	Ž.				PAGE	- - -	
YEAR	1985	1986	1967	1988	1989	1990	1661	1992	1993	1994	
	50.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	
PRODUCTION, HH UNITS/YEAR JET FUEL (BBLS)	14.56	29.11	29.11	29.11	29.11	29.11	29.11	29.11	29.11	29.11	
AMONIA (ST.) SULFUR (LT.)	0.05	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	
PRODUCT PRICES: \$/UNIT JET FUEL \$/BBL	58.35	58.85	58.85	58,85	58.85	58.85	58.85	58.85	58,85	58.85	
APTONIA \$/ST SULFUR \$/LT	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	155.00	
PRODUCT VALUES SPYCYEAR JET FUEL APPORTA	856.58 7,97	1713.16	1713.16	1713.16	1713.16	1713.16	1713.16	1713.16 5.94	1713.16	1713.16	1
			:						;		

1713.16 5.94 1.71

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PRODUCT VALUES \$PHYYEAR
JET FUEL
APPONIA
SULFUR

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ECONOMIC EVALUATION MODEL RUN NUMBER 2 ALL & VALUES IN MILLIONS			USAF SHA PRODUCTION	usaf shale oil study Production suffary	À		JP8+DIESEL-15% ROI	PAGE
YEAR	1995	19%	1997	1998	1999	2000		
	100.00	100.00	100.00	100.00	100.00	100.00		
PRODUCTION, MH UNITS/YEAP JET FUEL (BBLS)	29.11	29.11	29.11	29.11	29.11	29.11		i
APPONIA (ST.)	0.10	0.10	0.10	0.10	0.10	0.10		
SULFUR (LT)	0.03	0.03	0.03	0.03	0.03	0.03		
PRODUCT PRICES: \$/UNIT JET FUEL \$/BBL	58.85	58.85	58.85	58.85	58.85	58.85		į.
APPONIA \$/ST	155.00	155.00	155.00	155.00	155.00	155.00		
SULFUR \$/LT	105.00	105.00	105.00	105.00	105.00	105.00		

Appendix B.5

בהת יוסדהו מדונדהת מתו הנוחס הסוס / מד נוחנד

	14:04 NOV
REPORT	
MAXIMUM JP-4 CASE STANDARD OPTIMIZATION REPORT	
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USAF SHALE OIL TO FUELS PHASE IV CASE 1 - HAXIMIN JP-4 JET FUEL

GROSS MARGIN SEPTIME

		DESCRIPTION OF THE PROPERTY OF	- Contract					
SALES HARKET		PRODUCT	QUANTITY LIHITS MAXIMUM HINIMUM	TS FIXED	QUANTITY SOLD	FRICE	TOTAL \$70AY	
< < <		AV TURBINE FUEL JP-4 SULFUR AMHYDRUS AFFONIA			62563,75 BBL 67.79 TON 275.84 TON	-56.8200 95.4500 155.0000	56.8200.4692408£7 95.4500 8379.53 155.0000 42754.63	
TOTA	TOTAL SALES REVENJE	*					+743542.83	:
RAH HATERIAL PURCHASED	PURCHASED	MATERIAL	QUANTITY LIMITS		QUANTITY PURCHASED	PRICE	TOTAL	
		OCCIDENTAL SHALE OIL COLD TREATED WATER 50# STRIPPING STEAM		90000	90000.00 BBL 14418.41 BBL 159.45 TGN	\$0.000 .0245	0 3600000.00 5 353.25 10 .00	•
TOTAL	TOTAL RAW MATERIAL COST	1500					34.00361 08	

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - MAXIMLM JR-4 JET FUEL

OPERATING COST SUPPARY

PROCESS UNIT	HOOF	A TONE TO THE TOTAL TO THE TOTAL TOT			:			1	
•	2	CONTROL STREAM	-	3	QUANTITY	8	OPERATING COST	TSC	
				8879	10/10	\$/88L	*/10N	9/1	
FEED PREPARATION	COMPTNED MODES			,					
P. HYDDOTOFATING		_	_	900006	14441.92	.029556	18419	2660 05	
2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		COMPOSITE COST	-	90000	14441.92	204147	00110		
SUT INCHORUS TO THE	COMBINED MODES	COMPOSITE COST	-	00670	10.002.00		00000	69.67.69	
HIDROCHACKING	COMBINED MODES				13.0464	*04067	1.53820	22134.8	
NAPHTHA SPLITTER		_	- I	8400I.3	12408.39	.687944	4.69045	58200.93	
DEPARTANT ZED			<u> </u>	24850.2	2843.75	. 052302	45704	1200 71	
		COMPOSITE COST	-	17915.8	1966.07	OFEEDT	20107		
TOCK GAS INCAICH	COMBINED MODES	COMPOSITE COST	.		1116.1		60000	. 5T.C/11	
SULFUR PLANT	COMBINED MODES		. }		00.000		. 56680	756.88	
HYDROGEN PLANT			- 1		46.22		19.67483	1932.48	
SOLID LIATED TORATTAIN			: -		737, 30		19 50067	20101 72	
THE PART OF THE	COTETNED MODES	COMPOSITE COST	j	17516 2	1040 00	******			
ATTINE MEGENERATION	COMBINED MODES					****	1.189	3643.86	
APPONIA PLANT	COMPTNED MODES		- 1		78.55		17.81405	1749.71	
FIEL OTI STABILITICS	CONTRACT INDES		-		275.84		S ARRAI	741 47	
L SI ABILITER	COMBINED MODES	COMPOSITE COST		11824 2	2027 50	74.0		10.14	
CUTBINED FACILITIES	COMBINED MODES				75.1707	47CU/0	4/175	28.75	 1
		1		90000.0	14441.92	620860.	.61090	8622.57	
TOTAL PROCESS UNIT OPERATING COST	VIING COST								
							-	159772.79	
UTILITY PURCHASES U	UTILITY	THE PARTY OF THE P		!					
		ŝ	UNITS/D	*ZWIT	9,				
4	BOILER HATER	MLBS	4810 A1		01.76				
				2000	74.479				
TOTAL UTILITY PURCHASES				1	!				 i
,								14.4142	
UTILITY PRODUCTION COSTS	. עזרזזז	LNIT	UNITS/D	SOURCE	3C€		2		
	ACOR STEAM	3	, , ,						
			4564.99	REFINERY FUELS	r FUELS	12	1112.49		
TOTAL UTILITY PRODUCTION COSTS	COSTS								
	:						•	3112.49	
				1	-				!
OIAL UPERALING LUST							-	94 301341	
							•	44.60666	

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STANDARD OPTIMIZATION REPORTS, PS1.0 USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - MAXIMIM JP-4 JET FUEL

UNIVERSAL OIL PRODUCTS CO.

CAPITAL COST SUPPARY

PROCESS UNIT	700E	CONTROL STREAM	80 50	QUANTITY CO.D	198/1	CAPITAL COST	57	
FEED PREPARATION	COMBINED MODES		0.0000	14441 02	204976	6000		
L. P. HYDROTREATING	COMBINED MODES	COMPOSITE COST	90000	14441.92	i	1 944.91	E4 000 70	
HYDOOGBACKENING	COMBINED MODES		92629.0	14390.21		42362	90440.70	
NADIALA SOLUTION	COMBINED MODES	_	84601.3	12408.39		11.02282	16.1602 21	
DEBUTANT SED	COMBINED MODES		24850.2	2843.75		78742	22.246.04	
FUEL GAS TOFATED	COMBINED MODES	COMPOSITE COST	17915.8	1946.07	. 218830	2.01459	3920.52	
Sul Flat Plant	COTESINED FIGURES			1335.36		1.18287	1579.56	
HYDROGEN PI ANT	COMBINED FIGURES			98.22		57.37486	5635.41	
SOLD HATED TOFATTING	COMPTNED FIGURES		:	737,30		162,24250	11%20.79	
AMINE DEGENERATION	COMPTNED PROFES		17534.2	3069.99	. 530932	3.03242	9309.49	· ·
APPICIAL PLANT	COMPINED MODES			98.22		26.64155	2616.75	
FUEL OIL STABILIZED	Constitute Foots	_		275.84		1.50765	415.86	
COMBINED FACTUATIES	CONSTRED FIGURES		13824.2	2027.59		1.44524	2930.36	
	COLLEGE COLLEGE	COMPOSITE COST	90000.0	14441.92	. 115235	.71813	10371.11	
TOTAL							04 3116	

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USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JET FUEL MATERIAL BALANCE SUPPARY

SP.GS.		Š	TONS/D	HT%	HSCFD
.91650	90000.0	86.192 13.808	14441.92 2524.45 159.45	84.328 14.741 .931	
	104418.4	418.4 100.000	17125.61	100.000	
85	8870	Ķ	10NS/D	HTX	HSCFD
76135	92583.6	79.089	79.089 11297.75 87.79 275.84	65.969 .513 1.611	
	82583.8	79.089	79.089_11661.37	68.092	
SP.GR.	. 881/0	ΓΛX	TONS/D	71	HSCFD
.86400	0 13159.9	12.603	1990.75	11.624	
	13159.9	12.603	1990.75	11.624	
SP.GR.	88170	IV.	10NS/D	X LA	HSCFD
7.00000	15927.1	15.253	27,02 10.43 647.65 2788.60	. 159 . 061 3. 782 16. 283	
	15927.1	15.253	3473.69	20.283	
	111670.7	106.945	1670.7 106.945 17125.81	100.000	

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - MAXIMM JP-4 JET FUEL

		P. P.	PLANT CAPACITY SUFFIARY	INVESTMENT	INVESTMENT CAPACITY UNITS/DAY	TS/DAY			
רווח	CAPACITY	ITY CATEGORY	HAXIHUM	HINIHON	FIXED	LP INPUT	ACTUAL	INVESTMENT	
FEED PREPARATION	CAPACITY					90000	90004.9	9030000	
L.P. HYDROTREATING	CAPACITY					90000	6.40006	59430000	1
H.P. HYDROTREATING	CAPACITY					92630.0	92635.6	96390000	
HYDROCRACKING	CAPACITY					84600.6	84600.6	168517609	
NAPHTHA SPLITTER	CAPACITY					24851.3	24851.3	2335209	
DEBUT ANI ZER	CAPACITY				•	17916.8	17916.8	4088542	
FUEL GAS TREATER	CAPACITY	10NS/CD				1355.4	1335.4	1647258	
SULFUR PLANT	CAPACITY					98.5	8.8	5876921	
HYDROGEN PLANT	CAPACITY					737.3	737.3	124747396	
SOUR MATER TREATING	CAPACITY					17624.7	17624.7	9408468	
AHINE REGENERATION	CAPACITY					98.5	88.5	2728900	
APPOSTA PLANT	CAPACITY					275.8	275.8	433687	
FUEL OIL STABILIZER	CAPACITY		•			13824.1	13824.1	3055945	
COMBINED FACILITIES	CAPACITY					90000.0	90006	10815000	

TOTAL OPTIMIZED INVESTMENT

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIHIZATION REPORTS, PS1.0

	Ě	PRODUCT BLEN	BLENDING								
AV TURBINE FUEL JP-4	8879	TONS/D	SPGRa60F	HTZ SULF	VLZ AROM	SHOKE PT	FREEZ PT	D+L 293F	D+L 374F	D+L 473F	
HC KEROSINE: 300-520F	65556.5	9207.64	.8022	.0003	9 -6000	27.0000	-58,0000		38.0000	88.0000	
SPLITTER BOTTOMS	6390.3	852,56	. 7620	.0003	7.0000		-120,0000	98.0000	110.0000	100.000	
LT NAPHTHA JP-4	10637.0	1237.55	9999.	.0003	5.0000		-160.0000		120.0000	110.0000	
BLEND	62563.8	11297.75	.7814	.0003	7.9400	28.4000	-75.9353	25.0240	54.1331	91.7622	
SPECIFICATIONS		HAX	.7507	4000	25.0000	20.0000	-72.0000	20.0000	50.0000	30.000	
	ă	PRODUCT BLEN	BLENDING				:		i		
AV TURBINE FUEL JP-4	881/0	TONS/D	0+L 518F								
HC KEROSINE: 300-520F SPLITTER BOTTONS LT NAPHTHA JP-4	65556.5 6390.3 10637.0	9207,64 852.36 1237,55	100.0000		· !		<u>.</u>		:		İ
BLEND	65283.6	11297.75	100.0000				•				:
SPECIFICATIONS		HAX	100.0000								
	E .	PRODUCT BLEN	BLENDING	•							
SULFUR	BB(70	10NS/D	SPGR360F								
SULFUR		87.79				:	*		1		
BLEND		87.79						٠			
NO SPECIFICATIONS	:					I				To a second of manual of second	
	æ	PRODUCT BLEN	BLENDING								
ANHYDRUS AFTONIA	88/2	TONS/0	SPGR260F								ı
ANHYORUS ANHONIA		275.84									
BLEND	*	275.84		1			:				:
NO SPECIFICATIONS											

VERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0	STANDARD OPTI	HIZATION !	REPORTS, PS1.0	16:04 NOV 20, 81 PAGE 8
USAF SHALE OIL TO	FUELS			
CASE 1 - MAXIMUM JP-4 JET FUEL	JP-4 JET FUEL			The second secon
	BLE	NO FOR 41	BLEND FOR UTILITY PRODUCTION	
INERY FUELS	88.70	TONS/0	TONS/D SPGRa60F	
F.O. STAB BOTTOMS	13159.9	1990.75	.8640	
BLEND	13159.9	1990.75	0498.	
NO SPECIFICATIONS				
	And the state of t			

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - HAXIMLM JP-4 JET FUEL

DETAILED MATERIAL BALANCE FEED PREPARATION

16:04 NOV 20, 61 PAGE 10		the second secon			۵						-					0						
		:			MSCFD		90 404.90			29028.66						MSCFD			tr ata			
		,			1	93.47	3.24	100.00		4.62	21.42	69.16	8	100.00		MTX.	100.00	100.001	9	68.43	100.00	
				· •	TONS/D	12408.39	429.70	13274.99		613.10	2843.75	9207.64	6.07	13274.99		TONS/D	2843.75	2843.75	4K 11	1946.07	2843.75	
P51.0			33	:	Š	97.18	2.82	100.00			28.55 1 95	75.30		107.80		3	100.00	100.00		72.10	97.81	
ON REPORTS, P			MATERIAL BALANCE	YIELDS	8879	64601.3	2454.2	87055.5			24850.2 3460.8	65556.9		93847.5	MATERIAL BALANCE PLITTER : YIELDS	588	24850.2	24950.2		17915.8	24306.1	
STANDARD OPTIMIZATION	FUELS	IP-4 JET FUEL	DETAILED MATER	COMPOSITE YI	SP, GR.	. 6377	1.0000				98.69.	B022			DETAILED MAT NAPHTHA SPLI COMPOSITE YI	SP. GR.	.6536			.6204	i	
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV	CASE 1 - MAXIMUM JP-4 JET FUEL			CHARGE	SEPARATOR LIQUID(HP)	COLD TREATED HATER 50# STRIPPING STEAN	CHARGE	PRODUCTS	LP FLASH GAS(JP-4)	SOUR HATER (HC)	HC KEROSINE: 300-520F	SSOT	PRODUCTS		. SABOR	PROD FRACT OV'0: JP-4	CHARGE	PRODUCTS SPLITTER GAS; JP-6	SPLIT OV LIQ: JP-4 SPLITTER BOTTOMS	PRODUCTS	

MSCFD 2187.43

TONS/D

SP. Ga.

98.22 100.00 98.22 100.00 89.38 10.62

67.79 10.43

PRODUCTS SULFUR SULFUR PLANT LOSS

PRODUCTS

CHARGE HYDROGEN SULFIDE 100.00

98.25

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UNIVERSAL OIL PRODUCTS CO.	STANDARD OPTIMIZATION REPORTS, PS1.0	REPORTS, F	51.0			
USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMUM JP-4 JET FUEL	FUELS JP-4 JET FUEL					
	DETAILED MATERIAL BALANCE DEBUTANIZER COMPOSITE YIELDS	TERIAL BAL	NCE.			
	SP.GR.	8670	Š	TONS/D	MTZ	MSCFD
SPLIT OV LIQ:JP-4	.6204	17915.8	100.00	1946.07	100.00	
CHARGE		17915.6	100.00	1946.07	100.00	
PRODUCTS DEBUT OVER'D=JP-4 LT NAPHTHA JP-4	5464	6693.0	37.36 62.64	640.29	32.90 67.10	9451.01
PRODUCTS		17916.3	100.00	1946.07	100.00	
	DETAILED MATERIAL BALANCE FUEL GAS TREATER COMPOSITE YIELDS	TERIAL BAU EATER TELDS		Í	; ;	
A DO	SP.6R.	88/70	Ľ	TONS/D	H	MSCFD
LP FLASH GAS(JP-4)				613.10	45.91	29028.66
DEBUT OVER'D=JP-4 SPLITTER GAS:JP-4 F O STAB OWED:D	. 5464	6693.0	100.00	640.29 45.13	47.95 3.38	9451.01
CHARGE		6693.0	100.00	1335.36	100.00	0.5.000
PRODUCTS		:				•
HYDROGEN SULFIDE				18.65	1.40	415.25
4-40-00-01 TAI				635.04	45.40	9117.27
		!	,	44.28	3.32	896,15
			ļ	34.23	2.56	1960.49
PRODUCTS				1335.36	100.00	
	DETAILED MATERIAL BALANCE SULFUR PLANT COMPOSITE YIELDS	TERIAL BALVI T TELDS	PNCE			:

Tribaction National Planuere Corporate Value lue Corporate Value Corporate Value Corporate Value	UNIVERSAL OIL PRODUCTS CO. STANDARD OP- USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JET FUEL	STANDARD OPTIMIZATION REPORTS, P51.0 UELS >-4 JET FUEL	REPORTS, P	51.0			16:04 NOV 20, 81 PAGE	M .
SP.GR. BBL/O LVX TONS/O HTX 461.04 41.54 44.28 564.4 100.00 68.22 4.93 564.4 100.00 68.22 4.93 564.4 100.00 1384.94 100.00 DETAILED HATERIAL BALANCE SOLR HATER TREATING COPPOSITE YIELDS SP.GR. BBL/O LVX TONS/O HTX 1.0000 1558.4 20.29 1.0000 1558.4 20.29 1.0000 1558.4 20.29 1.0000 15643.6 60.06 1.0000 15643.6 90.36 555 1.0000 15643.6 90.36 555 1.0000 DETAILED HATERIAL BALANCE SOLR HATERIAL BALANCE SOLR HATERIAL BALANCE SOLR HATERIAL BALANCE SOLR HATERIAL BALANCE TONS/O 1069.99 100.00 96.22 100.00 96.22 100.00 96.22 100.00	!	DETAILED HAT HYDROGEN PLA COMPOSITE YI		3 ×				
## 663.04 43.54 65.64 100.00 643.27 45.64 645.00 1364.94 100.00 643.22 4.93 2.47 100.00 643.22 4.93 2.47 100.00 643.22 4.93 2.47 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 1364.94 100.00 100		SP. SP.	8870	r.	TONS/D	H	HSCFD	
### 1.0000 1564.4 100.00 68.22 4.93 ### 1.0000 1558.4 100.00 1394.94 100.00 DETAILED MATERIAL BALANCE SOLR HATER TREATING COPPOSITE YIELDS SP.GR. BBL/D LV: TONS/D HT? 1.0000 1558.4 20.29 623.02 20.29 I.0000 15643.4 20.34 2773.98 90.34 BR 1.0000 15643.4 90.34 2773.98 90.34 SP.GR. BBL/D LV: TONS/D HT; SP.GR. BBL/D LV: TONS/D HT; SP.GR. BBL/D LV: TONS/D HT; 98.22 100.00 98.22 100.00 98.22 100.00	FLASH: JP-4 OV'D=JP-4 II 6AS=JP-4 STAR OV: D	:	:		635.17	43.54 45.86 3.20	26804.15 9337.27 896.15	į
### 1.0000 1584.6 100.00 1364.94 100.00 DETAILED NATERIAL BALANCE SOUR HATER TREATING COCHOSTIE YIELDS SP.GR. BBL/D LVX TONS/D HTX 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 20.29 1.0000 1589.4 20.29 6.23.02 1.554.2 100.00 3669.99 100.00 DETAILED MATERIAL BALANCE ANTHE REGENERATION COCHOSTIE YIELDS SP.GR. BBL/D LVX TONS/D HTX SP.GR. BBL/D LVX TONS/D HTX 98.22 100.00 21 98.22 100.00 21	THA JP-4	5999.	\$96.4	100.00	68.22	4.93		
### 137.30 51.24 22% 647.65 46.76 1364.94 100.00 DETAILED NATERIAL BALANCE SOUR HATER TREATING COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX 1.0000 1558.4 20.29 623.02 20.29 1.0000 10535.1 60.09 1644.54 60.09 1.0000 10535.1 60.09 1644.54 60.09 1.0000 15643.6 90.36 602.43 19.62 1.0000 15643.6 90.36 2773.98 90.36 SP.GR. BBL/D LVX TONS/D HTX COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX 99.22 100.00 99.22 100.00 99.22 100.00	ARGE		586.4	100.00	1384.94	100.00		
DETAILED HATER TREATING SOUR HATER TREATING COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX 1,0000 1358-4 20.29 623.02 20.29 .HT 1,0000 1358-1 60.08 1644-54 60.08 1,0000 1440.8 19.62 602.43 19.62 17534.2 100.00 3649.99 100.00 DETAILED HATERIAL BALANCE AHINE REGENERATION COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX SP.GR. BBL/D LVX TONS/D HTX 98.22 100.00 21 98.22 100.00 21	N (97%) D.PLT.		:		737.30	53.24 46.76	22%23.91	:
BBL/D LV: TGNS/D HT: SP.GR. BBL/D LV: TGNS/D HT: 1.0000 1558-4 20.29 623.02 20.29 1.0000 1558-4 20.29 623.02 20.29 1.0000 1558-4 19.62 602.43 19.62 1.0000 1558-4 20.00 19.62 602.43 19.62 1.0000 1584-6 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 2773.98 90.36 29.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.40 29.52 100.00 200.00000000000000000000000000000	coucts			i	1384.94	100.00		
SP.GR. BBL/D LVX TONS/D HTX. 1.0000 1558.4 20.29 623.02 20.29 1.0000 10535.1 60.08 1844.54 60.08 1.0000 15534.2 100.00 3069.99 100.00 ER 1.0000 15843.4 90.34 2773.98 90.34 290.46 9.46 5.55 .18 DETAILED HATERIAL BALANCE AHINE RECENERATION COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX 98.22 100.00 21 98.22 100.00 21		DETAILED HAT SOUR HATER T	ERIAL BALA REATING ELDS	PCE.	• •			
HT 1.0000 3558.4 20.29 623.02 20.29 1.0000 10535.1 60.06 1644.54 60.08 1.0000 3440.8 19.62 602.43 19.62 17534.2 100.00 3069.99 100.00 15843.6 90.36 2773.98 90.36 200.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.46 290.46 9.40 290.22 100.00 200.00		S. S.	8870	r _V ;	10NS/D	774	HSCFD	
.4 90.36 2773.98 90.36 .6 90.36 2773.98 90.36 .6 90.36 3069.99 100.00 MLANCE D LVX TONS/D MTX 98.22 100.00 98.22 100.00 23	WATER - L.P.HT WATER - M.P.HT WATER (MC)	1.0000	3558.4 10535.1 3440.8	20.29 60.08 19.62	623.02 1844.54 602.43	20.29 60.08 19.62		
ER 1.0000 15843.6 90.36 2773.98 90.36 9.46 9.46 9.46 9.46 9.46 9.46 9.46 9.4	ARGE		17534.2	100.00	3069.99	100.00		
15843.6 90.36 3069.99 100.00 DETAILED MATERIAL BALANCE AHINE REGENERATION COMPOSITE YIELDS SP.GR. BBL/D LV% TONS/D MT% 98.22 100.00 98.22 100.00 98.22 100.00	SOUR HATER	3.0000	15843.6	90.36	2773.98 290.46 5.55	90.36	-	
DETAILED MATERIAL BALANCE AMINE REGENERATION COMPOSITE YAELDS SP.GR. BBL/D LV.: TONS/D MT.: 98.22 100.00 21 98.22 100.00 21	COUCTS	!		90.36	3069.99	100.00		
SP.GR. BBL/D LV% TONS/D MT% 96.22 100.00 96.22 100.00 21	•	DETAILED HAT AHINE REGENE COMPOSITE YI	ERIAL BALA RATION ELDS	SCE .	; 1	; ;		
96.22 100.00 96.22 100.00 96.22 100.00 96.22 100.00		SP. GB.	288	[%	TONS/D	HTX	HSCFD	
96.22 100.00	N SULFIDE		ţ		98.22	100.00	2187.43	:
96.22 100.00	ARGE				96.22	100.00		
98.22	N SULFIDE		-	! :	96.22	100.00	2187.43	
	DOUCTS				98.22	100.00		

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NOTE OF THE POSTS, BIANDARD OPTIMIZATION REPORTS, PSI.O		·
ANDARD	E3	CASE 1 - MANTHON 10 A 16T ELLE
	USAF SHALE OIL TO FUELS PHASE IV	9
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	DETAILED HA	HATERIAL BALANCE	ANCE			
•	APPONIA PLA	YIELDS		1	i i i	
CHARGE	SP.GR.	88/0	Ľ.	TONS/D	MT.X	HSCFD
AHONIA				290.46	100.00	
CHARGE				290.46	100.00	
PRODUCTS TREATED SOUR MATER ANHYDRUS APPIONIA	1.0000	63.5		14.62	5.03	
PRODUCTS		63.5	:	290.46	100.00	
;	DETAILED MATERIAN FUEL DIL STABILIA COMPOSITE YIELDS	MATERIAL BALANCE STABILIZER YIELDS	SACE.	; 1		
CHABGE	SP. GR.	88.70	Ľ	TONS/0	7	HSCFD
SEPARATOR LIQUID(HP)	. 6377	13824.2	100.00	2027.59	100.00	
CHARGE		13824.2	100.00	2027.59	100.00	
PRODUCTS F.O. STAB OVER'D F.O. STAB BOTTOMS	0,98	13159.9	95.19	36.64	1.82	2016.90
PRODUCTS	! !	13159.9	95.19	2027.59	100.00	The company of the state of the
:	DETAILED MA COMBINED FA	MATERIAL BALANCE FACILITIES YIELDS				
MCCOATC	SP.GR.	88170	ž.	TONS/D	X X	HSCFD
OCCIDENTAL SHALE OIL	. 9165	90000.0	100.00	14441.92	100.00	
CHARGE		90000.0	100.00	14441.92	100.00	
PRODUCTS OCCIDENTAL SHALE OIL	. 9165	0.00006	100.00	14441.92	100.00	
PR00UCTS		90000.0	100.00	19441.92	100,00	

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USAF SHALE OIL TO FUELS Phase IV CASE 1 - MAXIMIM JP-4 JET FUEL

	•	DETAILED USAGE OF LECTRIC POMER LCOSTS ARE ALLOCATE	DETAILED USAGE OF AN UNPOOLED UTILITY ELECTBIC POWER UNIT IS KUM COSTS ARE ALLOCATED TO PROCESS UNITS	PRICE IS	.04500/LNIT			
PROCESS UNIT	HODE	CONTROL STREAM	8870	TONS/D	UNIT/88L	UNITATON	UNITS/0	
L.P. HYDROTREATION L.P. HYDROTREATING H.P. HYDROTREATING H.P. HYDROCRACKING NAPHTHA SPLITTER DEBUTANIZER FUEL GAS TREATER SULFUR PLANT HYDROGEN PLANT HYDROGEN PLANT ANTHE REGENERATION	COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	90000.0 90000.0 92629.0 84601.3 24850.2 17915.8	14441.92 14441.92 14506.21 12408.39 2843.75 1946.07 1335.36 99.22 737.30	. 1080 1.9070 1.2677 7.0969 . 3911 . 1399	673 111.884 6.160 6.160 3.418 1.288 1.288 21.974 235.698	9720.85 117627.72 117422.70 600404.58 9719.95 2506.53 2158.30 173779.45	437 7723 5284 27018 437 112 7820 298
ATTONIA PLANT FUEL OIL STABILIZER COMBINED FACILITIES		COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	13824.2	98.22 275.84 2027.59 14441.92	. 1638	1.116	4004.76 915.86 2263.80	190

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CO. STANDARD OPTIMIZATION REPORTS, P51.0	
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FRSAL OIL	USA

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USAF SMALE UIL IO FUELS PHASE IV	4-90
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CASE 1 MANAGEMENT A SECTION OF THE PROPERTY OF
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		COOLING WATER LINIT IS HEAL. COSTS ARE ALLOCATED TO PROCESS UNITS	OCESS UNITS	PRICE 15	PRICE IS 03000/UNIT				
PROCESS UNIT	HODE	CONTROL STREAM	88/20	TONS/D	UNIT/BBL	WIT/TON	UNITS/D	Q.	
FEED PREPARATION	COMBINED HODES	COMPOSITE USAGE	90000.0	14441 92	:			:	
L.P. HYDROTREATING		COMPOSITE USAGE	90000.0	14441.92	9100	010	145.84	87.4	
H.P. HYDROTREATING		COMPOSITE USAGE	92629.0	14390.21		<u>:</u>		}	
HIDROCRACKING	COMBINED MODES	COMPOSITE USAGE	84601.3	12408.39	0850		3365,15	100.95	
NAPHTHA SPLITTER		COMPOSITE USAGE	24850.2	2843.75	.0299	. 261	742.22	22.27	
DEBUT WIZER		COMPOSITE USAGE	17915.8	1946.07	.0116	106	207.26	6.22	
FUEL GAS TREATER	COMBINED MODES	COMPOSITE USAGE		1335.36		•			
SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE		99.22					
HYDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		737.30		820	611.33	18.36	
SOUR MATER TREATING	COMBINED MODES	COMPOSITE USAGE	17534.2	3069.99	.1336	763	2342.76	70.28	
AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.22		32, 389	3181.27	95. 46	
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.84		3	1021.44	00.00	
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	13824.2	2027.59	.0200	117	277.17	A. 12	
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	900000	14441.92	1	; ;		!	

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - MAXIMLM JP-4 JET FUEL

	4	DETAILED USAGE OF A POOLED UTILITY REFINERY FUELS UNIT IS H-BTU COSTS ARE ALLOCATED TO UTILITY POOL	JLED UTILITY S HEBTU UTILITY POOL						
PROCESS UNIT	MODE	CONTROL STREAM	88170	10NS/D	UNIT/861.	UNITATON	UNITS/D	2	
FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING	COMBINED MODES COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92	.0218	.136	1%1.21		
HYDROCRACKING	COMBINED MOCES	COMPOSITE USAGE	92629.0	14390.21	.0482	. 310 971.1	4466.72		;
DEBUTANI ZER	COMBINED MODES	COMPOSITE USAGE	24850.2 17915.8	2843.75	.0640	. 559	1591.08		
FUEL GAS TREATER SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE	•	1335.36					
SOUR HATER TREATING	COMBINED MODES	COMPOSITE USAGE	17534.2	737.30	•	57.519	42408.30		
ANTINE REGENERATION ANTONIA PLANT	COMBINED MODES	COMPOSITE USAGE	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	98.22					
ruel oil STABILIZER COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	13824.2 90000.0	2027.59	.0484	. 331	672.15		:
TOTALS			:		:		65725.22		
AHOUNT PRODUCED FROM F.O. STAB BOTTOMS	F.O. STAB BOTTONS						69675.21		
APOLNT CONVERTED TO 600# STEAM	SOOB STEAM						4149 99		

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VERSAL .	-

COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES	COSTS ARE ALLOCATED TO UTILITY POOL		,	; ;			
HODE CO-BINED CO-BINED CO-BINED CO-BINED CO-BINED CO-BINED		UTILITY POOL			 - !		
CO-BINED CO-BINED CO-BINED CO-BINED CO-BINED CO-BINED	CONTROL STREAM	BB/O	TONS/D	UNIT/88L	UNIT/TON	UNITS/D	\$
COPBINED COPBINED COPBINED COPBINED	COMPOSITE USAGE	900006	14441.92			•	i I
COMBINED COMBINED COMBINED	COMPOSITE USAGE	90000.0	14441.92				
COMBINED	COMPOSITE USAGE	92629.0	14390.21				
COMBINED	COMPOSITE USAGE	84601.3	12408.39	:			-
	COMPOSITE USAGE	24850.2	2843.75				
COMBINED	COMPOSITE USAGE	17915.8	1946.07				
FUEL GAS TREATER COMBINED MODES	COMPOSITE USAGE		1335.36				
COMBINED	COMPOSITE USAGE		98.22		-7.593	-745.80	1
COMBINED	COMPOSITE USAGE		737.30				
COMBINED	COMPOSITE USAGE	17534.2	3069.99	. 2633	1.504	4616.45	
COMBINED	COMPOSITE USAGE		98.22		1.660	163.07	
COMBINED	COMPOSITE USAGE		275.84				
COMBINED	COMPOSITE USAGE	13824.2	2027.59				
COMBINED FACILITIES COMBINED MODES	COMPOSITE USAGE	90000	14441.92				
TOTALS		:				4033.71	
AHOUNT PRODUCED FROM 600# STEAM						4033.71	

509.78

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USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMIM JP-4 JET FUEL

916.07 -968.82 **CNITS/0** 824.35 120-2 9.327 UNIT/TON .424 -9.864 .0440 UNIT/88L 14441.92 14490.21 12408.39 12408.39 1844.07 1335.36 1346.07 1335.36 135.36 135.36 135.36 135.36 135.36 135.36 135.36 136.26 136. 10NS/D 13824.2 90000.0 90000.0 92629.0 84601.3-24850.2 17915.8 DETAILED USAGE OF A POOLED UTILITY
50% STEAM
COSTS ARE ALLOCATED TO UTILITY POOL 8679 17534.2 COMPOSITE USAGE
COMPOSITE USAGE
COMPOSITE USAGE
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COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE USAGE USAGE CONTROL STREAM COMPOSITE U
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COMPOSITE U MODES MODES MODES MODES MODES HOOES HOOES MODES CO-BINED 1 CO-BINED 2 CO-BINED 2 CO-BINED 2 CO-BINED 3 COMBINED COMBINED COMBINED H.P. HYDROTREATING
HYDROCRACKING
NAPHTHA SPLITTER
DEBUTANIZER
FUEL GAS TREATER
SULFUR PLANT
HYDROGEN PLANT
SOUR WATER TREATING
AHINE REGENERATION ATTONIA PLANT FUEL OIL STABILIZER COMBINED FACILITIES HYDROTREATING **PREPARATION** PROCESS UNIT FEED

TOTALS

AMOUNT PRODUCED FROM 600# STEAM

2419.91

4619.61

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USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JET FUEL

	:	DETAILED USAGE OF A POOLED UTILITY BOILER HATER UNIT IS HIBS	UTILITY					!
		COSIS AME ALLOCATED TO UTILITY POOL	LITY POOL					
PROCESS UNIT	HOOE	CONTROL STREAM	88/0	10NS/D	UNIT/BBL	UNITATON	UNITS/D	2
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	0.0000	14441 92				
L.P. HYDROTREATING	COPBINED MODES	COMPOSITE USAGE	90000	14441.92				
H.P. HYDROTREATING	COMBINED MODES		92629.0	14390.21				
HYDROCRACKING	COMBINED MODES		84601.3	12406.39	.0112	970.	946.76	ļ
NAPHTHA SPLITTER	COMBINED MODES		24650.2	2843.75			!	
DEBUTANIZER	COMBINED MODES	COMPOSITE USAGE	17915.8	1946.07				
FUEL GAS TREATER	COMBINED MODES	COMPOSITE USAGE		1335.36				
SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE		53.55			1860.89	
HIDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		737.30	•	16.687	12450.88	
SOUR MATER TREATI'S	COMBINED MODES	COMPOSITE USAGE	17534.2	3069.99				
AMINE REGENERATI.	COMBINED MODES	COMPOSITE USAGE		98.22				
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE	1	275.84				
FUEL OIL STABILIZER	COMBINED MODES		13824.2	2027.59				
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	0.00006	14441.92				
TOTALS							15258.54	!

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ATOUNT PRODUCED FROM CONDENSATE

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UNIVERSAL OIL PRODUCTS CO. STANDARO OPTIMIZATION REPORTS, PS1.0 USAF SHALE OIL TO FUELS PHASE IV CASE I - MAXIMUM JP-4 JET FUEL

	9/•		
	UNITS/D	-1733.03 -1462.05 -636.55 -824.35 -67.14 -4616.45 -1079.15	
	UNITATON	120 102 103 424 684 -1. 504 -10. 967	
	UNIT/BBL	0193 0158 0075 0460	
	TONS/D	14441.92 14441.92 14390.21 12408.39 2843.75 1946.07 1335.36 98.22 737.30 98.22 275.84	14441.92
D UTILITY PLBS TILITY POOL	88170	90000.0 92629.0 92650.3 24650.3 17915.8 17534.2	A0000.0
DETAILED USAGE OF A POOLED UTILITY CONDENSATE UNIT IS HLBS COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM		CONFUSTIE USAGE
ı	HODE	COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES	כשמיני השונים הח
	PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING NAPHTHA SPLITTER DEEUTANIZER FUEL GAS TREATER SOUEW PLANT HYDROGEN PLANT HYDROGEN PLANT ATTING ATTING ATTING ATTING ATTING FUEL OIL STABLIZER CUMMINEN	

TOTALS

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AMOUNT CONVERTED TO BOILER MATER

UNIVERSAL OIL PRODUCTS CO.	TS CO. STANDARD	OPTIMIZATION REPORTS, P51.0	1.0			16:04	16:04 NOV 20, 81	PAGE 22
USAF SHALE PHASE IV CASE 1 - M	USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMJM JP-4 JET FUE	JE L		,		The same same same same same same same sam	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	! ! !
	:	DETAILED USAGE OF AN UNPOOLED UTILITY CAT. & CHEMICALS UNIT IS \$ COSTS ARE ALLOCATED TO PROCESS UNITS	POOLED UTILITY S \$ PROCESS UNITS		PRICE IS 1.00000/UNIT			
PROCESS UNIT	HOOE	CONTROL STREAM	88/VB	TONS/D	UNIT/BBL	UNITATION	UNITS/D	9
FEED PREPARATION	COMBINED HODES	COMPOSITE USAGE	0.00006	14441.92	:			<u>;</u>
L.P. HYDROTREATING		COMPOSITE USAGE	900006	14441.92	.1145	.714	10308.64	10308.64
H.P. HYDROTREATING		_	92629.0	14390.21	.0417	. 268	3858.02	3858.02
HYDROCRACKING	COMBINED MODES	COMPOSITE USAGE	84601.3	12408.39	1820	809	6180.62	6180.62
NAPHTHA SPLITTER		_	24850.2	2843.75				
DEBUT ANI ZER		_	17915.8	1946.07				
FUEL GAS TREATER		COMPOSITE USAGE		1335.36				
SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE	1	98.22	3			
HIDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		737.30		5.004	3689.72	3689.72
SOUR HATER TREATING	COMBINED MODES	COMPOSITE USAGE	17534.2	3069.99				
AMINE REGENERATION		COMPOSITE USAGE		98.22		. 283	27.76	27.76
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.84	;			
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	13824.2	2027.59				
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000	14441.92				
TOTALS		1		1	!		24064.75	24064.75

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USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JET FUEL

PROCESS UNIT UTILITY CONSUMPTION SUPPARY

				:
CONDENSA TE MLBS	-1733.0	-824.4	-4616.4	-10418.7
BOILER W ATER MLBS	946.B	1860.9		15258.5
50# STEA M MLBS	-261.8	824.4 -968.8	916.1	509.8
150# STE AH MLBS	•	-745.8	4616.4	4033.7
REFINERY FUELS MMBTU	1961.2 4466.7 14625.8	42408.3	672.1	65725.2
COOLING MATER MGAL	145.9	207.3	2342.8 3181.3 3023.4 277.2	136%.5
600# STE AN MLBS	1733.0 1462.0 636.6	21.6	!	21.5
ELECTRIC POWER KUM	9720.9 171627.7 117422.7 600404.6	2506.5 2158.3 173779.5	6640.4 4004.8 935.9 2263.8	1101184.9
HOCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING NAPHTHA SPLITTER	DEBUTANIZER SULFUR PLANT HYDROGEN PLANT	SUCH HAIEN INEATING ATINE REGENERATION ATTONIA PLANT FUEL OIL STABILIZER	TOTAL CONSUMPTION

							Appei	ndix	B.5	(Cont.)
16:04 NDV 20, 81 PAGE 24									· · · · · · · · · · · · · · · · · · ·	
CO. STANDARD OPTIMIZATION REPORTS, P51.0	USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JET FUEL	PROCESS UNIT UTILITY CONSUMPTION SUFFARY	CAT. & C HEMICALS	10308.6	6180.6	3689.7		24064.7		
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV CASE 1 - MAXIMLM JP-4 JI		PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING		HYDROGEN PLANT SOUR WATER TREATING	ATTONE MESENERATION APPOINTA PLANT FUEL OIL STABILIZER	TOTAL CONSUMPTION	-231-	

Appendix B.6

UNIVERSAL OIL PRODUCTS CO. STA	STANDARD OPTIMIZATION REPORTS, PS1.0	51.0		16:04 NOV 20, 61	1. 61 PAGE
PHASE IN CASE 2 - JP-4 JET FUEL	TPUEL + DF-2/0FH DIESEL			1	*
	GROSS MARGIN SLIFTARY				
SALES MARKET	PRODUCT	QUANTITY LIMITS	QUANTITY SOLD	33 100	TOTAL
444	AV TURBINE FUEL JP-9 DF-2/DF-H DIESEL SULFUR AMYDRUS AFFONIA	·	48559.86 EBL 33184.22 BBL 67.76 TGL	58,8200 58.5200 95,4500	2856. 1951.
TOTAL SALES REVENUE		1	51 64-679	155.0000	42691.06
RAH MATERIAL PURCHASED	MATERIAL	QUANTITY LIMITS	QUANTITY FURCHASED) PRICE	4859254.15 Total
	OCCIDENTAL SHALE OIL COLD TREATED HATER SOR STRIPPING STEAN	0.00006	90000.00 BBL 14342.57 BBL 154.52 TON	.0245	3600000.00 351.39
GROSS MARGIN	COST	· · · · · · · · · · · · · · · · · · ·			3600351.39
	;				JZ 206 8571

190491.12.

TOTAL OPERATING COST

P51.0
ION REPORTS,
STANDARD OPTIMIZATION
CO. STANDA
DIL PRODUCTS
UNIVERSAL 0)

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USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

OPERATING COST SUPPARY

PROCESS UNIT	70F	CONTROL STREAM	!	8	QUANTITY	7	OPERATING COST	OST	
				88 /2	TONO	198/ +	*/10V	2	
FEED PREPARATION	COMBINED MODES	COMPOSITE COST		90000.0	14441.92	.029556	18419	2660.05	:
L.P. HYDROTREATING	COMBINED MODES	_		90000.0	14441.92	. 294147	1.83308	26473.23	
H.P. HYDROTREATING	COMBINED MODES	COMPOSITE COST		92629.0	14390.21	. 238964	1.53820	22134.96	
HYDROCRACKING	COMBINED MODES	_		81987.1	12024.97	.618072	4.21406	50673.94	
NAPHTHA SPLITTER	COMBINED MODES	٦	:	11463.6	1292.32	.083930	74321	960.46	
DEBUTANI ZER	COMBINED MODES	U		8380.3	901.49	. 113850	1.05835	954.09	
FUEL GAS TREATER	COMBINED MODES	_			778.09		.91020	708.21	
SULFUR PLANT	COMBINED MODES	_			98.18		19.67483	1931.73	
HYDROGEN PLANT	COMBINED MODES	COMPOSITE COST			421.42		43.68431	18409.54	
SOUR WATER TREATING	COMBINED MODES	_		17427.9	3051.37	. 208795	1.19253	3638.86	
AHINE REGENERATION	COMBINED MODES	_			98.18		17.81405	1749.04	
AMMONIA PLANT	COMBINED MODES	COMPOSITE COST			275.43		2.68881	740.57	
FUEL OIL STABILIZER	COMBINED MODES		;	16438.4	2411.00	.067122	45764	1103.38	
PARTIAL OXIDATION	COMBINED MODES	_			285.42		162.49114	46377.54	
COMBINED FACILITIES	COMBINED MODES	COMPOSITE COST		90000.0	14441.92	.098029	.61090	8822.57	
TOTAL PROCESS UNIT OPERATING COST	ERATING COST	· · · · · · · · · · · · · · · · · · ·		!				167336.18	
UTILITY PURCHASES	עזותא	CNI	UNITS/D	\$/WIT	2				
	BOILER HATER	HLBS	6310.09	20000	3155.05				:
TOTAL UTILITY PURCHASES	ES							3155.05	
UTILITY SALES	UTILITY	דאט	UNITS/D	#/UNIT	4.				:
	600# STEAM 50# STEAM	MLBS MLBS	422.80 623.15	.00010	 				
TOTAL UTILITY SALES	1					:		or.	

UNIVERSAL DIL PRODUCTS CD. STANDARD OPTIMIZATION REPORTS, P51.0 USAF SHALE DIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

CAPITAL COST SUPPARY

PROCESS UNIT	HODE	CONTROL STREAM		QUANTITY		CAPITAL CO	
			88179	10/0	188/t	NOT/	2.
FEED PREPARATION	COMBINED MODES	TOCH BITTE COST	00000	20.663		1	
O HYDROTORATING				36.4555		24400	2074, 20
DITIKUK DEDIKAT KATA	_	_	90000.0	14441.92		3.94621	56990.78
A.P. HIUNDIREATING	_	_	0.0420	14190 21		4 4034.9	46.700
HYDROCBACKTAG	•			****		34634.0	10.46474
CONTRACTOR OF THE PARTY OF THE	_	Τ.	81987.1	12024.97		13.13572	157956.60
MAINTAN ANIELS	_		11443.6	1292, 32		1.17586	1510 60
DEBUTANI ZER	_		I USIA	07 (00	1000	01.70	
FUEL GAS TOFATED	•					6.4/434	76.1007
	_	_		776.09		1.54%1	1205.73
	_	_		98.18		57, 38256	5611.99
HIDROGEN PLANT	_	_		421.42		178 9271	75401 99
SOUR HATER TREATING	COMBINED MODES	COMPOSITE COST	17427 0	105 17	E19913	14010	20.000
AMINE REGENERATION	_					3.03.40	17.6/24
APPOINT A DI ANT	•	_		. ÷		56.64666	2616.25
	_					1.50877	415.56
TOEL OIL STABILIZER	_		16438.4		194388	1, 32515	1105.41
PARTIAL OXIDATION	_	COMPOSITE COST	;			10107 101	100656 07
COMBINED FACILITIES	-		00000			1000	10.404.47
	•	•	40000	74 - 7 + 4 + 7	115235	. 71813	10371.11

16:04 NOV 20, 81 PAGE 5			HSCFD			HSCFD			ISCFD			HSCFD			
			HTX	84.418 14.679 .903	100.000	MTX	39.034 28.287 .513	69.444	MTX	6.342	8.342	HTX	. 156 . 061 1. 976 16. 194 3. 827	22.215	100.000
			10NS/0	14441.92 2511.17 154.52	17107.61	10NS/0	6677.79 4839.21 87.76 275.43	11680.18	TONS/D	1427.04	1427.04	TONS/0	26.76 10.43 338.05 2770.40 654.74	3800.39	17107.61
P51.0		¥RY	- -	86.254 13.746	100.000	Ľ	46.539 31.803	78.342	N'S	9.041	150.9	Ľ	15.165	15.165	102.548
		ANCE SUPPL	8870	90000.0 14342.6	104342.6	88	48559.9 33184.2	81744.1	86.5	9433.5	9433.5	867	15823.2	15823.2	107000.7
STANDARD OPTIMIZATION REPORTS,	USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL	HATERIAL BALANCE SURHARY	SP. GB.	, 91650		SP. GR.	,78543 . 83290		59 SP. GR.	. 86400		SP. GR.	1.00000		
UNIVERSAL OIL PRODUCTS CO.	USAF SMALE OIL TO FUELS PMASE IV CASE 2 - JP-4 JET FUEL		CHARGE	OCCIDENTAL SHALE OIL COLD TREATED WATER 50% STRIPPING STEAM	TOTAL CHARGE	PRODUCTS	AV TURBINE FUEL JP-4 DF-2/DF-M DIESEL SULFUR ANYDRUS ANYONIA	TOTAL PRODUCTS SOLD	STREAMS CONVERTED TO UTILITIES	F.O. STAB BOTTOMS	TOTAL STREAMS CONVERTED	STREAMS NOT UTILIZED	LOSS SULFUR PLANT LOSS CO2HYO.PLT. TREATED SOUR HATER PARTIAL OXIDATLOSS	TOTAL NOT UTILIZED	TOTAL PRODUCTS HADE

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

		PLANT	PLANT CAPACITY SUPPARY	INVESTMENT CAPACITY UNITS/DAY	APACITY UNI	TS/DAY			
UNIT	CAPACITY	CATEGORY	HAXIHUH	HINIMCA	FIXED	LP INPUT	ACTUAL	INVESTHENT	İ
FEED PREPARATION	CAPACITY	BBLS/CD				90000	90004.9	9030000	
L.P. HYDROTREATING	CAPACITY	BBLS/CO				0.00006	6,40006	59430000	•
H.P. HYDROTREATING	CAPACITY	B815/CD				92630.0	92635.6	96390000	
HIDROCRACKING	CAPACITY	BBLS/CO				81986.5	81986.5	164726166	
NAPHTHA SPLITTER	CAPACITY	BBLS/CO				11444.7	11444.7	1584723	
DEBUTANIZER	CAPACITY	Bals/CD				8380.8	8380.8	2796286	!
FUEL GAS TREATER	CAPACITY	TONS/CD				778.1	778.1	1257405	
SULFUR PLANT	CAPACITY	TONS/CO				98.2	98.5	5875452	
HYDROGEN PLANT	CAPACITY	TONS/CD				421.4	421.4	78635478	
SOUR WATER TREATING	CAPACITY	88LS/C0				17518.3	17518.3	9673242	
AMINE REGENERATION	CAPACITY	TONS/CD				98.5	88.8	2728375	
APPONIA PLANT	CAPACITY	TONS/CD				275.4	275.4	433365	
FUEL OIL STABILIZER	CAPACITY	BBLS/CO				16438.2	16438.2	3332381	
PARTIAL OXIDATION	CAPACITY	TONS/CD	:	:		205.4	285.4	114145893	,
COMBINED FACILITIES	CAPACITY	BBLS/CD				90000.0	90004.9	10815000	

IVERSAL UIL PRUUUCIS CO.	STANDARD OPT	IMIZATION	OPTIMIZATION REPORTS, PS1.0	51.0				16:04	16:04 NOV 20, 81	PAGE	
USAF SHALE OIL TO FUELS PHASE IV	FUELS										
CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL	FUEL + DF-2/DI	FM DIESEL	!								•
	ă	PRODUCT BLENDING	DING								
TURBINE FUEL JP-4	88	TONS/D	SPGR360F	TONS/D SPGRa60F HTZ SULF	VL% AROH		SMOKE PT FREEZ PT	D+L 293F	D+L 374F	D+L 473F	:
HC KEROSINE:300-520F SPLITTER BOTTOMS LT NAPHTHA JP-4 DH	40776.0 2742.7 5041.1	5727.13 365.92 584.74	.8022 .7620	.0003	9.0000 7.0000 2.0000	35.0000	27.0000 -58.0000 35.0000 -120,0000 35.0000 -160,0000	2.5000 98.0000 120.0000		98.0000 100.0000	
BLEND	48559.9	66.77.39	. 7854	.0003	8.1600	28.2800	28.2800 -72.0907	20.0919	50.5793	90.9617	
SPECIFICATIONS		MAX	. 7507	. 4000	25.0000	20.0000	-72.0000	20.0000	50.0000	90.0000	i
	Ě	PRODUCT BLENDING	SNIO			:	:				
TURBINE FUEL JP-4	BB/70	10NS/D	TONS/D D+L 518F								
HC KEROSINE:300-520F SPLITTER BOTTOMS LT NAPHTHA JP-4 DH	40776.0 2742.7 5041.1	\$727.13 365.92 584.74	5727.13_100.0000 365.92_100.0000 584.74_100.0000	!							1
BLEND	48559.9	6677.79	100.0000								
SPECIFICATIONS		HAX	100.0000						!	·	:

UNIVERSAL OIL PRODUCTS CO.	STANDARD OPTIMIZATION REPORTS, P51.0	HIZATION	REPORTS, P	51.0				16:04	16:04 NOV 20, 81	PAGE	o
USAF SHALE CIL TO FUELS	FUELS										
CASE 2 - JP-4 JET FUEL + DF-2/DFH DIESEL	FUEL + DF-2/DF	H DIESEL	i.					! !		•	
	6 4	PRODUCT BLENDING	DING								
DF-2/DF-M DIESEL	BBL/D	TONS/D	SPGR260F	WTZ SULF	FLASH PT	CETANE I	D+L 700F	VIS 100F	POUR PT	כנסעם פי	,
HC DIESEL:520-700F	33184.2	4839.21	.8329	. 0003	268.0000	56.0000	100.0000	16.2400	0000	5.0000	
BLEND	33184.2	4839.21	.8329	. 0003	268.0000	56.0000	100.0000	18.2400	0000	5.0000	
SPECIFICATIONS	:	HAX	.8607	. 7000	133.0000	45.0000	100.0000	18.4100	0000	5.0000	:
	æ	PRODUCT BLENDING	DING								
DF-2/DF-M DIESEL	88/78	TONS/D	0+1, 675F	D+L 725F				1	i		
HC DIESEL:520-700F	33184.2	4839.21	90.0000	100.0000							
BLEND	33184.2	4819.21	90,0000	100.000	1	: !					;
SPECIFICATIONS		HAX	90.0000	100.0000							
	æ	PRODUCT BLENDING	DING						:	!	
SULFUR	8870	TONS/D	SPGRa60F								
SULFUR		87.76			i						
BLEND		87.76									
NO SPECIFICATIONS	•		:				i :		1	:	
	P8	PRODUCT BLENDING	DING					•			
ANHYDRUS APPONIA	88.7	TONS/D	SPGR360F	!	:	:					
ANHYDRUS APPONIA		275.43									
BLEND		275.43					:	; ; !	1	i	· ·
NO SPECIFICATIONS											
	ane .	BLEND FOR UT	FOR UTILITY PRODUCTION	UCTION	4						
REFINERY FUELS	8872	TONS/D	SPGRa60F								
F.O. STAB BOTTOMS	9433.5	1427.04	.8640		,			•			
BLENO	9433.5	1427.04	.8640								
NO SPECIFICATIONS				:							

16:04 NOV 20, 81 PAGE 9				MSCFD						HSCFD		77.76906			1/72.1B			MSCFD		72.706.00			
		:	i	X X	100.00	100.00	100.00	100.00	;	MTX	95.66	3.13	100.00	4.13	. \$4. 95.32 .03	100.00	•	MX	88.34	8	100.001	11.32 88.62 .05	100.00
	į.	ŧ	!			14441.92	14441.92	14441,92	:	TONS/D	14441.92	472.83	15097.29	623.02	79.57 14390.21 4.48	15097.29		10NS/0	14390.21	1621.92	16289,43	1844.54 14435.97 8.92	16289.43
P51.0			:			100.00	100.00	100.00		Š	60.76	2.91	100.00	3.84	36.95	103.76	Ä	[%	16.06	60.6	100.00	10.34 96.60	106.94
ON REPORTS, PS		MATERIAL BALANCE	<u> </u>	88C9	90000.0	90000.0	90000.0	90000	MATERIAL BALANCE OTREATING YIELDS	88.70	90000.0	2700.6	92700.6	3558.4	92629.0	96187,4	MATERIAL BALANCE OTREATING : YIELDS	88170	92629.0	9263.6	9.566101	10535.1 -98425.5	108%0.6
STANDARD OPTIMIZATION R	.E OIL TO FUELS JP-4 JET FUEL + DF-2/DFH DIESEL	DETAILED MATERIAL FEED PREPARATION		SP. GS	. 9165		. 9165		DETAILED MATERIAL I L.P. HYDROTREATING COMPOSITE YIELDS	SP. GR.	5916`	1.0000		1.0000	. 6673		DETAILED MATERIAL (M.P. HYDROTREATING COMPOSITE YIELDS	SP. GR.	18873	1.0000	-	1.0000	
UNIVERSAL OIL PRODUCTS CO. STAM	USAF SMALE OIL TO FUELS PMASE IV CASE 2 - JP-4 JET FUEL			CHARGE	OCCIDENTAL SHALE OIL	(ARGE	PRODUCTS DEASMED SMALE OIL	PRODUCTS			DEASHED SHALE DIL		CHARGE	PRODUCTS SOUR HATER - L.P.HT	SEPARATOR LIQUID(LP) LOSS	PRODUCTS		0 0 0 1	SEPARATOR LIQUID(LP)	COLD TREATED HATER	CHARGE	PRODUCTS SOUR MATER - H.P.HT SEPARATOR LIQUID(HP) LOSS	PRODUCTS

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USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

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; ; [MSCFD		*6.63V01		24226.60	 		!		:	: ! !	MSCFD				580.80	
	H	93.63	3.24	100.00	3.06	10.06	4.55 87.55	44.54	8.8.	100.001	i	H	100.00	100.001		1.93 69.76 28.31	100.00
	TONS/D	12024.97	416.42	12642.91	192.62	1292.32	583.81	4819 21	7.82	12842.91		10NS/0	1292.32	1292.32		24.92 901.49 365.92	1292.32
	ראג	97.18	2.82	100.00		13.56	3.95	3 2		105.18	. SC	Š	100.00	100.00		73.23	97.20
ILED HATERIAL BALANCE ROCRACKING POSITE YIELDS	68/79	81987.1	2378.4	84365.5		11443.6	3334.4	40776.0		88738.3	TERIAL BALLITTER	88/0	11443.6	11443.6		6380.3	11123.0
DETAILED MATERIAN HYDROCRACKING COMPOSITE YIELDS	SP.GR.	. 6377	3.0000			0979.	1.0000	2208	550.		DETAILED MATERIAL BALANCE NAPHTHA SPLITTER COMPOSITE YIELDS	SP.GR.	. 6450			. 4144	
		SEPARATOR LIQUID(HP)	HYDROGEN (972) COLD TREATED WATER 50# STRIPPING STEAM	CHARGE	PRODUCTS I B EI ASH GAS: IB-4-DH	PROD FRACT OV: JP4-DH	SOUR HATER (HC)	HC KEROSINE: 300-520F	LOSS LOSEL: 350-700F	PRODUCTS	· •		CHARGE PROD FRACT OV: JP4-DH	CHARGE	BBONICTS	SPLITTER GAS=JP4-DH SPLIT OV LIQ=JP4-DH SPLITTER BOTTOHS	PRODUCTS

UNIVERSAL OIL PRODUCTS CO. STANDA	STANDARD OPTIMIZATION REPORTS, P51.0	REPORTS, P	51.0				16:04 NOV 20, 81 PAGE 11
USAF SHALE OIL TO FUELS PHASE IV							
JP-4 JET FUEL	+ OF-2/OFM DIESEL						
	DETAILED MATERIAL DEBUTANIZER COMPOSITE YIELDS	ERIAL BALANCE	Z CE	r C			
00 A M O	SP.GR.	8870	L	TONS/D	HTX	MSCFD	
SPLIT OV LIQ=JP4-DM	.6144	8380.3	100.00	901.49	100.00		
CHARGE		6380.3	100.00	901.49	100.00		
PRODUCTS DEBUT OVER'D=JP4-DH LT NAPHTHA JP-4 DM	.5418	3339.0	39.84 60.15	316.75 5 84.74	35.14 64.86	4802.32	
PRODUCTS		8380.2	100.00	901.49	100.00		i de la companya de l
	DETAILED MATERIAL BALANCE FUEL GAS IREATER COMPOSITE YIELDS	MATERIAL BALA IREATER TIELDS	I NCE		;	;	
u () () () () () () () () () () () () ()	SP. GR.	8870	rv.	TONS/0	MTX	MSCFD	
CALANCE DEBUT OVER'D=JP4-DH SPLITTER GAS=JP4-DH F.O. STAB OVER'D	.5418	3339.0	100.00	392.62 316.75 24.92 43.81	50.46 40.71 3.20 5.63	24226.60 4802.32 580.80 2400.68	
CHARGE		3339.0	100.00	778.09	100.00		
PRODUCTS HYDROGEN SULFIDE TRT LP FLASH: JP4-DH TRT DEB OV'D=JP4-DH TRT SPLIT GAS=JP4-DH TRT F.O. STAB OV'D		•		18.61 582.92 311.77 24.09 40.70	2.39 49.21 40.07 3.10	414.41 23998.91 4691.33 562.38 2333.22	
PRODUCTS				778.09	100.00		
	DETAILED MATERIAL BALANCE SULFUR PLANT COMPOSITE YIELDS	ERIAL BALA ELDS	NCE.		:		
T A A B B B B B B B B B B B B B B B B B	SP. GR.	BB170	I.K.	TONS/D	HTX	HSCFD	
HYDROGEN SULFIDE				98.18	100.00	2186.59	
CHARGE				98.18	100.00		
PRODUCTS SULFUR SULFUR PLANT LOSS	1	;		87.76 10.43	89.38 10.62		
PRODUCTS				98.18	100.00		

USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/OFM DIESEL

UNIVERSAL OIL PRODUCTS CO.

UNIVERSAL DIL PROCUCTS CO. STANDARD OPTIMIZATI USAF SMALE DIL TO FUELS PRASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIF	STANDARD OPTIMIZATION REPORTS, P51.0 UELS UEL + DF-2/DFM DIF DETAILED MATERIAL BALANCE	REPORTS, PS	2 W			; ·	16:04 NOV 20, 81 PAGE 13
	COMPOSITE YIELDS	ELDS	8	10MS/0	ž	MSCFD	
CHARGE APPONIA		;		290.05	100.00		
CHARGE				290.05	100.00	;	
PRODUCTS TREATED SOUR WATER ANY DRUS ATTOWA	1.0000	83.4		14.60	5.03	,	
PRODUCTS		93.4		290.05	100.00		
	DETAILED MAY FUEL DIL STA COMPOSITE Y	ID MATERIAL BALANCE IL STABILIZER ITE YIELDS	33				
	Sp. GS	88 79	Š	TONS/D	MTX	MSCFO	
CHARGE SEPARATOR LIQUID(HP)	.6377	16438.4	100.00	2411.00	100.00	•	
CHARGE		16438.4	100.00	2411.00	100.00		
PRODUCTS F.O. STAB DVER'D F.O. STAB BOTTOMS	0,98.	15648.4	95.19	43.81	1.82 98.18	2400.68	
PRODUCTS		15648.4	95.19	2411.00	100.00		
	DETAILED MATERIAL BALANCE PARTIAL OXIDATION. COMPOSITE YIELDS) MATERIAL BALA OXIDATION FE YIELDS	NCE .	:	; ;		
	SP. GR.	88/0	Š	TONS/D	HTX	MSCFD	
CHARGE F.O. STAB BOTTOMS	0,98.	6215.0	100.00	940.16	100.001		The same of the sa
CHARGE		6215.0	100.00	940.16	100.00		
PRODUCTS H) DROGEN (97%) PARTIAL OXIDATLOSS				285.42 654.74	30.36	86889. 98	
PRODUCTS				940.16	100.00	•	

16:04 NOV 20, 81 PAGE 14			/D MTX MSCFD	14441.92 100.00	90000.0 100.00 14441.92 100.00	90000.0 100.00 14441.92 100.00	90000.0 100.00 1441.92 100.00
			TONS/D		14441.9	14441.9	14441.
51.0		PCE	ĽĶ	100.00	100.00	100.00	100.00
REPORTS, P		ERIAL BALA ILITIES ELDS	8870	900000	90000.0	165 90000.0	900000
STANDARD OPTIMIZATION REPORTS, P51.0	USAF SMALE OIL TO FUELS PMASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL	DETAILED MATERIAL BALANCE COMBINED FACILITIES COMPOSITE YIELDS	SP. GR.	. 9165		. 9165	
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL		u con in in in in in in in in in in in in in	OCCIDENTAL SHALE DIL	CHARGE	PRODUCTS OCCIDENTAL SHALE OIL	PRODUCTS

1513488.45

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0

USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

	:	DETAILED USAGE OF AN UNPOOLED UTILITY ELECTRIC POWER UNIT IS KUM COSTS ARE ALLOCATED TO PROCESS UNITS	OLED UTILITY KWH OCESS UNITS	PRICE IS	.04500/UNIT	1		i
PROCESS UNIT	HODE	CONTROL STREAM	88C0	TONS/D	UNIT/88L	UNIT/TON	UNITS/D	•
SEED DOFFDADATION	SHOW NEW TONCO	FORBOSTTE LISAGE	0.00006	14441.92	. 1080	.673	9720.85	437.
LEG TACTOR LEGG	COMPANED MODES	COMPOSITE USAGE	90000.0	14441.92	1.9070	11.884	171627.72	7723.
H.P. HYDROTREATING	COMBINED MODES		92629.0	14390.21	1.2677	8.160	117422.70	5284.
HIDBOCRACKING	COMBINED MODES		81987.1	12024.97	6.1047	41.622	500503.29	22522
NAPHTHA SPLITTER	COMBINED MODES		11443.6	1292.32	. 3912	3.464	4476.35	201.
DEBUT ANI ZER	COMBINED MODES		8380.3	901.49	. 1399	1.301	1172.47	52.
FUEL GAS TREATER	COMBINED MODES			778.09				
Sulf G Prant	COMBINED MODES			98.18		21.974	2157.47	
HYDROGEN PLANT	COMBINED MODES			421.42		238.910	100682.00	
SOLID HATER TREATING	COMBINED MODES		17427.9	3051.37	. 3787	2.163	6600.30	
AMINE REGENERATION	COMBINED MODES			98, 18		40.773	4003.22	
ACHOUTA DI ANT	COMBINED HODES			275.43		1.191	24.42	i
FIEL OIL STABLIZER	COMBINED MODES	COMPOSITE USAGE	16438.4	2411.00	.1638	1.116	2691.89	121.
PARTIAL OXIDATION	COPBINED MODES			285.42		2072.400	591495.71	•••
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92				

P51.0	
STANDARD OPTIMIZATION REPORTS,	
UNIVERSAL DIL PRODUCTS CO.	

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USAF SHALE DIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFH DIESEL

		į	: :		
	0/4			,	ş
	UNITS/D	1733.03	21.57	-4020.37	-2908.89
	UNITATON	.120	. 220	-14.086	
	UNIT/BBL	.0193			:
,	TONS/D	14441.92 14441.92 14390.21	12024-77 1292-32 901-49 778-09 96,18 421-42 3051-37 275-43	2411.00 265.42 14441.92	;
LINI IS HUBS	8870	90000.0	81987.1 11443.6 8380.3 17427.9	16438.4	
DETAILED USAGE OF A POOLED UTILITY 600# STEAM LNIT IS MLBS COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	
	HODE	COMBINED MODES COMBINED MODES COMBINED MODES	COTBINED MODES COTBINED MODES COTBINED MODES COTBINED MODES COTBINED MODES COTBINED MODES COTBINED MODES		
	PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING	HIDROCRACKING NAPHTHA SPLITTER OEBUTAHIZER FUEL GAS TREATER SULFUR PLANT HIDROGEN PLANT SOUR WATER TREATING AMINE REGERATION	ATTINIA PLANI FUEL OIL STABILIZER PARTIAL OXIDATION COMBINED FACILITIES	TOTALS AMOUNT SOLD

AMOUNT CONVERTED TO 150# STEAM

40027.60

PAGE 17	· !		6		0 •	64.57	10.25	2.91			10.98	9.69	95.40	90.57	69.6	842.02	
16:04 NOV 20, 81	:	:	UNITS/D		73.00	2152.47	341.62	8.91			366.13	2328.61	3180.05	3018.95	329.58	28067.22	
N 50:91			UNIT/TON		010.	179	. 264	.107		:	.869	. 763	32.389	10.081	.137	96.338	
		.03000/UNIT	UNIT/BBL	. 7,00	9700.		.0299	.0116		:		. 1336		!	.0200		
		PRICE IS	TONS/D	14441.92	14390.21	12024.97	1292.32	901.49	776.09	98.18	421.42	3051.37	98.18	275.43	2411.00	285.42	14441.92
0.1		OOLED UTILITY S MGAL PROCESS UNITS	BB 170	90000 0	92629.0	61967.1	11443.6	6380.3				17427.9		•	16438.4		0.00006
OPTIMIZATION REPORTS, PS1.0	F-2/DFM DIESEL	DETAILED USAGE OF AN UNPOOLED UTILITY COOLING LATER . UNIT IS MGAL COSTS ARE ALLOCATED TO PROCESS UNITS	CONTROL STREAM	COMPOSITE USAGE	_	_	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE
STANDARD	ō		MODE	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL		PROCESS UNIT	FEED PREPARATION	ATING			DEBUTANIZER				(2)	ATION		æ		COMBINED FACILITIES

TOTALS

PORTS, P51.0		
UNIVERSAL OIL PRODUCTS CO. STANDARO OPTIMIZATION REPORTS, PS1.0		CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL
. STANDARD	TO FUELS	ET FUEL + OF
. PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV	SE 2 - JP-4 JI
UNIVERSAL OIL	USA	Ž

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FEED PREPARATION COMBINED MODES COMPOSITE USAGE L. P. HYDROTREATING H. P. HYDROTREATING H. P. HYDROTREATING H. P. HYDROTREATING H. P. HYDROTREATING H. P. HYDROTREATING H. P. HYDROTREATING COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE COMBINED MODES COMPOSITE USAGE ANTONIA PLANT COMBINED MODES COMPOSITE USAGE ANTONIA PLANT COMBINED MODES COMPOSITE USAGE C		OSTS ARE ALLOCATED TO UTILITY MOUL					
COPBINED MODES COMPOSITE LO CO		88170	10NS/D	UNIT/BBL	UNIT/TON	UNITS/D	۵/•
COPBINED MODES COMPOSITE L COPBINED MODES COMPOSITE L	HODES COMPOSITE	90000	14441.92	:			· · · · · · · · · · · · · · · · · · ·
COPBINED MODES COMPOSITE LOCABINED MODES COM	HODES COMPOSITE	90000	14441.92	.0218	.136	1961.21	
COMBINED HODES COMPOSITE L COMBINED HODES COMPOSITE L	MODES COMPOSITE	92629.0	14390.21	.0482	.310	4466.72	
COMBINED MODES COMPOSITE (COMBINED MODES COMPOS	HODES COMPOSITE	61967.1	12024.97	.1721	1.173	14110.10	:
COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPELNED MODES COMPOSITE (COPERNED MODES COPERNED MODES COMPOSITE (COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES	HODES COMPOSITE	11443.6	1292.32	0590.	.567	732.75	
COPELNED MODES COMPOSITE (COPERNED MODES COPERNED MODES COMPOSITE (COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED MODES COPERNED	HODES COMPOSITE	8380.3	901.49				
COPBINED MODES COMPOSITE (COPBINED MODES COPBINED MODES COMPOSITE (COPBINED MODES COMPOSITE (COPBINED MODES COPBINED MODES COMPOSITE (COPBINED MODES COPBIN	DOES COMPOSITE (778.09				
COPELNED MODES COMPOSITE LO COPELNED MODES COMPOSITE LO COPELNED MODES COMPOSITE LO COPENDED MODES COMPOSITE LO COPENDED MODES COMPOSITE LE COPENDED MODES COPENDED MODES COMPOSITE LE COPENDED MODES COMPOSITE LE COPENDED MODES COPENDED MODES COMPOSITE LE COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDED MODES COPENDE MODES COPE	COMPOSITE (98.18				
COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES COMPOSITE (COPENCE HODES HODES COMPOSITE (COPENCE HODES HODES COMPOSITE (COPENCE HODES HO	DOES COMPOSITE (421.42		26 .90¢	23980.61	
COPELNED MODES COMPOSITE I COPELNED MODES COMPOSITE I COPELNED MODES COMPOSITE I COPELNED MODES COMPOSITE I COPELNED MODES COMPOSITE I	COMPOSITE (17427.9	3051.37				
COMBINED MODES COMPOSITE I COMBINED MODES COMPOSITE I COMBINED MODES COMPOSITE I ES COMBINED MODES COMPOSITE I	COMPOSITE I		98.18				
ER COMBINED MODES COMPOSITE I COMBINED MODES COMPOSITE I ES COMBINED MODES COMPOSITE I	COMPOSITE I	,	275.43	: : : : : : : : : : : : : : : : : : : :			
COMBINED MODES COMPOSITE (DOES COMPOSITE	16438.4	2411.00	9840.	.331	799.25	
COMBINED MODES COMPOSITE	COMPOSITE I		285.42		14.149	4038.35	
	ODES COMPOSITE	90000.0	14441.92				
TOTALS	•	:	i	:	**	50088.99	
						00 0000	

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0	USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL
UNIVERSAL OIL PRODU	USAF SHALE PHASE IV CASE 2 - JI

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		DETAILED USAGE OF A POOLED UTILITY 150# STEAM. UNIT IS MLBS COSTS ARE ALLOCATED TO UTILITY POO	A POOLED UTILITY UNIT IS HLBS ED TO UTILITY POOL		1			1
PROCESS UNIT	HODE	CONTROL STREAM	8870	10NS/D	UNIT/BBL	UNITATON	UNITS/D	\$
FFFO DOFDADATION	STONE THE STONE							. !
	COLOTINED FILLES		90000.0	14441.92				
L. P. HIUNUINEALING	COMBINED MODES	COMPOSITE USAGE	90000	14441.92				
H. P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.0	14390.21				
HYDROCRACKING	COMBINED MODES	COMPOSITE USAGE	81987.1	12024.97	0185	-, 126	126 -1519.0K	
NAPHTHA SPLITTER	COMBINED MODES		11443.6	1292.32				!
DEBUTANI ZER	COMBINED MODES		8380.3	901.49				
FUEL GAS TREATER	COMBINED MODES			778.09				
SULFUR PLANT	COMBINED MODES	_		98.18		-7.591	1745.51	
HYDROGEN PLANT	COMBINED MODES			421.42			431911	:
SOUR HATER TREATING	COMBINED MODES		17427.9	3051.37	. 2633	1.504	6588.56	
AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.18		1.660	163.00	
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.43	,			
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	16438.4	2411.00				
PARTIAL OXIDATION	COMBINED MODES	COMPOSITE USAGE		285.42				
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92				
TOTALS	•	: :			:		2486.10	:

AMOUNT PRODUCED FROM 600# STEAM

8

623.15

P51.0
RD OPTIMIZATION REPORTS, P
. STANDA
L OIL PRODUCTS CO.
UNIVERSAL

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	USAF SHALE OIL TO FUELS	PHASE IV	CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

i								
	\$		1					1
	UNITS/D		-955.99	385.57	-968.45	915.71		
	UNIT/TON		e. 079	. 428	-9.864	9.327		;
	UNIT/BBL		0117	0940.				
	TONS/D	14441.92	12024.97	1292.32 901.49	778.09 98.18	421.42 3051.37 98.18	275.43 2411.00	285.42 14441.92
DF A POOLED UTILITY UNIT IS HLBS ATED TO UTILITY POOL	8879	90000.0	81987.1	11443.6 8380.3	٠	17427.9	16438.4	0.00000
DETAILED USAGE OF A POOLED UTILITY SOW STEAM COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM		COMPOSITE USAGE	COMPOSITE USAGE			COMPOSITE USAGE	COMPOSITE USAGE
	MODE	COMBINED MODES	COPBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES ::	COMBINED MODES
	PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING	H.P. HYDROTREATING HYDROCRACKING	NAPHTHA SPLITTER	FUEL GAS TREATER	HYDROGEN PLANT SOUR MATER TREATING	ATTO RECENERATION ANTONIA PLANT FUEL DIE STABILIZER	PARTIAL OXIDATION COMBINED FACILITIES

TOTALS

AMOUNT SOLD

6310.09

15625.15

P51.0	
STANDARD OPTIMIZATION REPORTS,	
UNIVERSAL OIL PRODUCTS CO. 9	

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USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL

	2																•
!	UNITS/D	1			1201.05				•	6974.54					3589.39		
	UNITATON				.266 1201.05				18.946	16.550					12.576		
	UNIT/88L				.0390												
	TONS/D	14441.92	14441.92	14390.51	12024.97	1292.32	901.49	778.09	98.18	451.45	3051.37	98.18	275.43	2411.00	265.42	14441.92	
WIT IS HLBS	88.7p	90000.0	0.00006	95629.0	81987.1	11443.6	6380.3				17427.9			16438.4		0.00006	
DETAILED USAGE OF A POOLED UTILITY BOILER WATER UNIT IS MLBS COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPUSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	
	HOOE	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	
	PROCESS UNIT	FEED PREPARATION	L.P. HIDROTREATING	H.P. HYDPOTREATING	HYDROCRACKING	NAPHTHA SPLITTER	DEBUT ANI ZER	FUEL GAS TREATER	SULFUR PLANT	HIDROGEN PLANT	SOUR WATER TREATING	AMINE REGENERATION	APPONIA PLANT	FUEL OIL STABILIZER	PARTIAL OXIDATION	COMBINED FACILITIES	

TOTALS

AMOUNT PRODUCED FROM CONDENSATE

AMOUNT PURCHASED

P51.0		
EPORTS,		
STANDARD OPTIMIZATION REPORTS, P51.0		-2/DFM DIESEL
STANDARD	FUELS	FUEL + DF.
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV	CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL
UNIVERSAL		

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		DETAILED USAGE OF A POULEU UILLITT CONDENSATE LINIT IS MLBS COSTS ARE ALLOCATED TO UTILITY POOL	S PLBS UTILITY POOL		i		; ;	:
PROCESS UNIT	MODE	CONTROL STREAM	88170	TONS/D	UNIT/BBL	UNIT/TON	UNITS/D	9,
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	0.00006	14441.92	ŧ			
L.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	0.00006	14441.92	0193	120	-1733.03	
H.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.0	14390.21	0158	102	-1468.05	
HIDROCRACKING	COMBINED MODES	COMPOSITE USAGE	A1987.1	12024.97		-		
NAPHTHA SPLITTER	COMBINED MODES	COMPOSITE USAGE	11443.6	1292.32				
DEBUT ANI ZER	COMBINED MODES	COMPOSITE USAGE	6360.3	901.49	0460	428	- 385.57	
FUEL GAS TREATER	COMBINED MODES	COMPOSITE USAGE		778.09				
SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE		98.18		684	-67.12	i
HYDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		421.42				
SOUR HATER TREATING	COMBINED MODES	COMPOSITE USAGE	17427.9	3051.37	2633	-1.504	-4588,56	
AHINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.18		-10.987	-1078.74	
APPONIA PLANT	COMBINED MODES	COMPOSITE, USAGE		275.43			1	
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	16438.4	2411.00				
PARTIAL OXIDATION	COMBINED MODES	COMPOSITE USAGE		282.45				
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92				

TOTALS

AMOUNT CONVERTED TO BOILER HATER

P51.0	
). STANDARD OPTIMIZATION REPORTS, PS1.0	
OPTIMIZATI	
STANDARD	FUELS
8	ار 10
PRODUCTS	USAF SHALE OIL TO FUELS
OI L	USA
UNIVERSAL OIL PRODUCTS CO.	

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	OF-2/DFM DIESE	
	•	
	4 JET FUEL	
	4-dC -	
, , , , ,	CASE 2 -	

		CAT & CHEMICALS INTO A CHOCKED UTILITY	OCEO UTILITY	21 20100	T100000 (31 30100			
		COSTS ARE ALLOCATED TO PROCESS UNITS	OCESS UNITS		5	i	:	: !
PROCESS UNIT	MODE	CONTROL STREAM	8870	TONS/D	UNIT/BBL	UNIT/TON	UNITS/D	•
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	900000	14441.92	i	!		
L.P. HYDROTREATING	COMBINED MODES	_	900000	14441.92	.1145	.714	10308.64	10308.64
H.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.0	14390.21	.0417	. 268	3858.02	3858.02
HIDROCRACKING	COMBINED MODES	COMPOSITE USAGE	81987.1	12024.97	0578	394	4742.65	4742.65
NAPHIHA SPLITTER	COMBINED MODES	_	11443.6	1292.32				
DEBUTANIZER	COMBINED MODES	COMPOSITE USAGE	9380.3	901.49				
FUEL GAS TREATER	COMBINED MODES	COMPOSITE USAGE		778.09				
SULFUR PLANT	COMBINED MODES	_		98.18				
HYDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		421.42	-	4.979	2098.39	2098.39
SOUR MATER TREATING	COMBINED MODES	COMPOSITE USAGE	17427.9	3051.37				
AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.18		. 283	27.75	27.75
AHHONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.43				
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE	16438.4	2411.00				
PARTIAL OXIDATION	COMBINED MODES	COMPOSITE USAGE		285.42		4.223	1205.31	1205.31
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000	14441.92				

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UNIVERSAL DIL PRODUCTS CO. STANDARD OPTIHIZATION REPORTS, PS1.0
USAF SHALE DIL TO FUELS
PHASE IV
CASE 2 - JP-4 JET FUEL + OF-2/DFH DIESEL

PROCESS UNIT UTILITY CONSUMPTION SUPPARY

PROCESS UNIT	ELECTRIC POWER KM	600# STE AM MLBS	COOLING HATER MGAL	REFINERY FUELS HABTU	150# STE AM MLBS	SO# STEA M MLBS	BOILER W ATER MLBS	CONDENSA TE MLBS
FEED PREPARATION	9720.9	6 1161	0 11 7	6		,		
1. P. HYDROTREATING	117422.7	1462.0	r.057	4466.7				-1/62.0
HDROCRACKING	500503.3		2152.5	14110.1	-1520.0	-956.0	3201.0	
NAPHTHA SPLITTER	4.97.4		341.8	732.7				•
DEBUTANI ZER	1172.5		8			385,6		-385.6
SULFUR PLANT	2157.5	21.6			-745.5	4.69.4	1860.2	-67.1
HYDROGEN PLANT	100682.0	-2105.2	366.1	23980.6			6974.5	
SOUR HATER TREATING	\$ 0099		2328.6		4588.6	•		-4588.6
MINE REGENERATION	4003.2		3160.1		163.0	915.7		-1078.7
MYONIA PLANT	934.5		3018.9					
FUEL OIL STABILIZER	2691.9	;	329.6	799.2				
PARTIAL OXIDATION	591495.7	-4050.4	28067.2	4038.3			3589.4	
TOTAL CONSUMPTION	1513488.5	-2908.9	40057.6	50089.0	2486.1	-623.2	15625.2	-9315.1

UNIVERSAL OIL PRODUCTS CO.	S CO. STANDARD OPTIMIZATION RE. CRTS, PS1.0	16:04 NOV 20, 81 PAGE 25
USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL	USAF SHALE OIL TO FUELS PHASE IV CASE 2 - JP-4 JET FUEL + DF-2/DFM DIESEL	
	PROCESS UNIT UTILITY CONSUMPTION SUPPARY	
PROCESS UNIT	CAT. & C HEMICALS	
FEED PREPARATION L.P. HIDROTREATING H.P. HYDROTREATING	10308.6 3858.0	
HIDHOCHACKING NAPHTHA SPLITTER DEBUTANIZER SUIFIR PLANT	4742.6	
HIDROGEN PLANT SOUR HATER TREATING	2098.4	
AHINE REGENERATION APPONIA PLANT ELIE OTI ETABLITED	7.73	
PARTIAL OXIDATION	1205.3	
TOTA_ CONSUMPTION	22240.7	

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USAF SHALE OIL TO FUELS PHASE IV) FUELS					
CASE 3 - MAXIMIN JP-8 JET FUEL	JP-B JET FUEL	•				
	GROSS MARGIN SUFFIARY	SUFFIARY				
SALES HARKET	PRODUCT	HAXINLH HININLH	S FIXED	QUANTITY SOLD	PRICE	TOTAL 6/0AY
< <	UNLEADED GASOLINE			7%3,79 BBL	58.800	4.8
•	AV IUMBINE PUEL JA-6 SULFUR			72667.97 BBL 87.77 TON	58.800	4272876.64
TOTAL CALES DESCRIPE	ATHURAS ATTURES		į	276.68 TON	155.0000	42665.11
IOIAL SALES MEVENUE	••		ŀ			4792410,45
MAM MATERIAL PURCHASED	MATERIAL	QUANTITY LIMITS	6 0	QUANTITY PURCHASED	FRICE	TOTAL
	OCCIDENTAL SHALE OIL COLD TREATED WATER 50% STRIPPING STEAM		90000.0	90000.00 BBL 14749.31 BBL 169.56 TON	0000	3600000.00
TOTAL RAW MATERIAL COST	COST					
GROSS MARGIN						3600361.36

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USAF SHALE OIL TO FUELS
PHASE IV
CASE 1 - MAXIMUM JP-8 JET FUEL

OPERATING COST SUTHARY

		: I	ppendix B.7	(Cont.)	
	:				
3ST 4/D	2660.05 26473.23 22134.96 60932.81 1251.84	1144.45 655.14 726.66 1932.09 29553.74 17253.34 4376.83	1749.36 743.93 980.52 8822.57	2760.43	2830.48
OPERATING COST \$/TON	1.6419 1.63306 1.53820 4.61775 47664	5.57700 .61770 19.67483 39.28601 2.21537 5.62530	17.81405 2.68881 .79034 .61090	.02	2830.48
0 \$788∟	. 029556 . 294147 . 238964 . 677281 . 054444	. 532947 . 532947 . 295563 . 750106	.115919		
QUANTITY D TON/D	14441,92 1441.92 14300.21 13195.34 2626,40	117.47 117.47 1176.39 93.20 752.27 778.81 778.42	98.20 276.68 1240.63 14441.92 84. 52	T \$/D 0 2780.43 SOURCE	REFINERY FUELS
B8	90000.0 90000.0 92629.0 89966.8 22993.1	1229.3 1229.3 5837.5 5837.6	8458.7 90000.0 840.6	\$/UNI	,
.	20 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	55 55 55 55 55 55 55 55 55 55 55 55 55	51 51 51 51	UNITS/D 5560.86 UNITS/D	4151.36
CONTROL STREAM	COHPOSITE COST COHPOSITE COST COHPOSITE COST COHPOSITE COST COHPOSITE COST		COMPOSITE COST COMPOSITE COST COMPOSITE COST COMPOSITE COST	MLBS	MLBS
100	COMBINED HODES COMBINED HODES COMBINED HODES COMBINED HODES COMBINED HODES		COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES ATING COST	WILITY BOILER HATER S UTILITY	600# STEAM
PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HIDROCRACKING NAPHTHA SPLITTER DEBUTANIZER	DEPROPAILZER FUEL GAS TREATER SULFUR PLANT HYDROSEN PLANT NAPHTHA HYDROTREATER UOP PLATFORHING SOUR WATER TREATING	AHINE REGENERATION CO-BINED APPONIA PLANT CO-BINED FUEL DIL STABILIZER CO-BINED CO-BINED FACILITIES CO-BINED OUTHY UNIT CO-BINED TOTAL PROCESS UNIT OPERATING COS	UTILITY PURCHASES B TOTAL UTILITY PURCHASES UTILITY PRODUCTION COSTS	60 TOTAL UTILITY PRODUCTION COSTS TOTAL OPERATING COST

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

CAPITAL COST SUPPLARY

PROCESS UNIT	HOOE	CONTROL STREAM)) ()	QUANTITY PRI ZD	ida/•	CAPITAL COST		
			3	5		5	7.	
FEED PREPARATION	COMBINED MODES	COMPOSITE COST	0.0000	14441 92		04004	02 0370	
L.P. HYDROTREATING	COMBINED MODES	_	00000	16661 00			DC . 7 C DO . 7	
H. P. HIDBOTDEATING	Condition of		0.000	74444		77056.6	30,70.78	
	COLDINGO FIORES	_	92629.0	14390.21		6.42342	92434.31	
A LONGCARCH ING	COMBINED MODES	COMPOSITE COST	80%6.8	13195.34		12.80446	168959.22	
NAPHINA SPLITTER	COPBINED MODES	COMPOSITE COST	22993.1	2626.40		.82011	2151.01	
DEBUTANIZER	COMBINED PODES	_	16633.9	1804.78	.227102	2.09310	3777.60	
DE PROPARII ZER	COMBINED MODES	_	1229.3	117.47		7.79117	915.24	
FUEL GAS TREATER	COMBINED MODES	~		1176.39		1.26026	1482,56	
SULFUR PLANT	COMBINED MODES	_		98.20		57.37868	5634.67	
HYDROGEN PLANT	COMBINED MODES	_		752.27		61.67252	121621.61	
NAPHTHA HYDROTREATER	COMBINED MODES	_	5837.5	778.81		4.19902	3270.22	
UOP PLATFORMING	COMBINED MODES		5837.6	778.42		13.29436	10346.56	
SOUR WATER TREATING	COMBINED MODES	Ξ.	17927.7	3139.68	526209	3.00544	9433.73	
AMINE REGENERATION	COMBINED MODES	COMPOSITE COST		98.20		26.64422	2616.49	
APPOSITA PLANT				276.68		1.50535	416.50	
FUEL DIL STABILIZER	COMBINED MODES	_	8458.7	1240.63		1.84761	2292.20	
COMBINED FACILITIES	COMBINED MODES	Ξ.	0.00006	14441.92	.115235	.71813	10371.11	
DOTHY CNIT	COFBINED MODES		9.058	84.52				

TOTAL

501378.11

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMLM JP-8 JET FUEL

MATERIAL BALANCE SURMARY

	300 10101.		5			
CHARGE	SP.GR.	88.70	ראי	TONS/D	MTX	HSCFD
OCCIDENTAL SHALE DIL COLD TREATED WATER 50% STRIPPING STEAM	. 91650	90000.0	85.919 14.081	14441.92 2582.39 169.56	83.995 15.019 .986	
TOTAL CHARGE		104749.3	100.000	17193.86	100.000	
PRODUCTS	SP. GB.	82	! !	TONS/D		HSCFD
UNIEADED GASOLINE AV TURBINE FUEL JP-8 SULFUR ANHYDRUS APHONIA	.80500	7963.8 72668.0	7.603	1041.78 10242.09 87.77 276.68	6.059 59.568 .510 1.609	App
TOTAL PRODUCTS SOLD		80631.8	76.976	11648.32	67.747	pend
STREAMS CONVERTED TO UTILITIES	SP. GA.	88/0	riv.	TONS/D	HTX.	1x B
F.O. STAB BOTTOMS LT NAPHTHA JP-8	.86400	8052.2 6918.6	7.687	1218.09	7.084	.7 ((
TOTAL STREAMS CONVERTED		14970.8	14.292	2022.78	11.765	Cont
STREAMS NOT UTILIZED	SP.GR.	68170	Š	TONS/D	нтх	USSFD
LOSS SULFUR PLANT LOSS Ç02HYD.PLT. TREATED SOUR HATER	1.00000	16315.7	15.576	27.54 10.43 	.160 .061 3.653 16.614	
TOTAL NOT UTILIZED		16315.7	15.576	3522.76	20.488	
TOTAL PRODUCTS HADE		111918.2	106.844	106.844 17193.86	100.000	

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

PLANT CAPACITY SUPPARY INVESTMENT CAPACITY UNITS/DAY

														•			
INVESTHENT	903000	59430000	00000196	176200124	2246246	1919493	954469	1546099	5876153	126811967	1610174	10792066	9818011	2728626	01717	2390435	10615000
ACTUAL	90004.9	90004.9	92635.6	89966.1	22993.9	16634.3	1229.2	1176.4	98.2	752.1	5837.5	5837.8	18018.5	8	276.7	8458.6	6.40006
LP INPUT	90000	90000	92630.0	89966.1	22993.9	16634.3	1229.2	1176.4	98.2	752.3	5937.5	5837.8	18018.5	98.2	276.7	9459.6	90000
FIXED						;											
HINIMGH																	
HAXIMUH			!														
CAPACITY CATEGORY	BBLS/CD	BBLS/CD	BBLS/CD	BBLS/CD	BBLS/CD	B81,5/C0	BBLS/CD	TONS/CD	TONS/CD	TONS/CD	88LS/CD	BBLS/CO	B6LS/CD	TO:45/CD	TONS/CD	BBLS/CO	BBLS/CD
CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY	CAPACITY
CNIT	FEED PREPARATION	L.P. HYDROTREATING	H.P. HYDROTREATING	HI DROCRACKING	NAPHTHA SPLITTER	DEBUTANI ZER	DE PROPANI ZER	FUEL GAS TREATER	SULFUR PLANT	HIDROGEN PLANT	NAPHTHA HYDROTREATER	UOP PLATFORMING	SOUR HATER TREATING	AMINE REGENERATION	APPIONIA PLANT	FUEL OIL STABILIZER	COMBINED FACILITIES

TOTAL OPTIMIZED INVESTMENT

Appendix B.7 (Cont.)

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

	æ	PRODUCT BLEN	ENDING								
UNLEADED GASOLINE	8870	TONS/D	TONS/D SPGRAGOF HTX SULF	HTX SULF	RES OCTN	HOT OCTN (R+H)12	. (R+H)12	RVP INDX VL% AROM D+L 131F	VL% AROM	D+L 131F	
LT NAPHTHA JP-8 100 RON PLATFORHATE HIXED BUTANES	2498.0 4625.2 6 40.6	290.54 666.71 84.52	.6643	.0003	75.0000 100.0000 98.5000	73.0000 68.6000 94.3000	74.0000 94.3000 96.4000	175.0000 55.1000 1280.0000	2.0000 72.6000 .0000	51.0000	
BLEND	7963.8	1041.78	1747.	.0003	92.0000	84.3084	88.1542	88.1542 222.0000	42.7920	27.1330	•
SPECIFICATIONS		HIN		.1000	92.0000	82.0000	87.0000	222.0000	55.0000	10.0000	
	Æ .	PRODUCT BLENDING	DING				1	1			1
UNLEADED GASOLINE	887	TONS/D	TONS/D D+L 171F D+L 235F	D+L 235F	D+L 365F	0+L 437F					
LT NAPHTHA JP-8 100 RON PLATFORMATE MIXED BUTANES	2498.0 4625.2 840.6		% 0000 7.0000 100.0000	100,0000 38,0000 100,0000	290.54 96.0000 100.0000 100.0000 666.71 7.0000 38.0000 100.0000 84.52 100.0000 100.0000 100.0000	100.0000					
BLEND	7963.8	1041.78	44, 7327	63.9916	63.9916 100.0000	100.0000					
SPECIFICATIONS		ÄH	50.0000	50.000	90.0000	100.0000					

Appendix B.7 (Cont.)

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - HAXIMUH JP-8 JET FUEL

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			1																					
	D+L 401F	40.0000	40.0000	10.0000																				
	FLASH PT	100.0000	100.0000	100.0000						1								:						
	VLY NAPH FLASH PT	1.5000	1.5000	3.0000											•			:				;		
	FREEZ PT	-58.0000	-58.0000	-58.0000								; ;		:										!
	SMOKE PT	26.0000	26.0000	20,0000				!				;						:						ı
	HTX SULF VLX AROH	10,0000	10.0000	25.0000								; ;				į				· •				:
	HTX SULF	.0003	.0003	.4000										:				:		9		;		
2		.8050	. 8050	.6398	DING	D+L 572F	100.0000	300,000	100.0000	DING	SPGR360F				DING	SPGR360F		!		RECYCLE STREAM BLENDING	SPGRa60F	.5743	.5743	
	BBL/D TONS/D SPGR860F	72668.0 10242.09	72668.0 10242.09	HAX	PRODUCT BLENDING	TONS/D D+L	72668.0 10242.09 100.0000	10242.09 100.0000	HAX	PRODUCT BLENDING	TONS/D	17.78	77.78		PRODUCT BLENDING	TONS/D SPGR	276.68	276.68		CYCLE STR	TONS/D SPGR	84.52	94.55	
	88.70	72668.0	72668.0	:	8	88/D	72668.0	72668.0			88/0				H.	88170				æ	88.70	840.6	9.05	1
	AV TURBINE FUEL JP-8	HC KEROSINE: 300-550F	BLEND	SPECIFICATIONS		AV TURBINE FUEL JP-8	HC KEROSINE: 300-550F	9LEND	SPECIFICATIONS		SULFUR	SULFUR	BLEND	NO SPECIFICATIONS		ANHYDRUS APPONIA	ANHYDRUS AHDNIA	BLEND	NO SPECIFICATIONS		MIXED BUTANES	MIXED C4'S:JP-8	BLEND	NO SPECIFICATIONS
										-26	2-	1				:								

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HALE OIL IV - MAXIM	USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL		:	į,			:		
		BLEND FOR UTILITY PRODUCTION	TY PRODUCTION						
	88.70 		SPGRa60F						
LT NAPHTHA UP-6	6918.6	1218.09 804.69	0 , 000 1,00	-		!			
BLEND NO SPECIFICATIONS	14970.8	2022.78	7177.						
							: ! !		
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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

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		The same of the sa					Арре	endi	ж І	3.7	(Con	t.)								
:	MSCFD	;					MSCFD	6. 6.0072	27.76006		9, 6,	7. Ot . 5/. 7			HSCFD		20 30 5 . 51		; ; ;	
	MTX	100.00	100.00	100.00	100.00		¥ 7.	95.66	3.13	100.00	4,13	.5. 95.32 .03	100.00		HT%	88.34	 8	100.00	11.32 86.62 .05	100.00
į	TONS/D	14441.92	14441.92	14441.92	14441.92		TONS/D	14441.92	472.83	15097, 29	623.02	14390.21	15097, 29	:	TONS/D	14390,21	1621.92	16289.43	1844.54 14435.97 8.92	16289.43
# 5	7.	100.001	100.00	100.00	100.00		LV %	97.09	2.91	100.00	3.84	99.92	103.76	3	ľ.	90.91	9.03	100.00	10.34	106.94
ERIAL BALANCE TION ELDS	88170	900006	90000	900000	0.00006	ERIAL BALA EATING ELDS	8879	0.00006	2700.6	92700.6	3558.4	92629.0	96187.4	ERIAL BALA EATING ELDS	88 7 8 7	92629,0	9263.6	101892.4	10535.1	108%0.6
DETAILED MATERIAL FEED PREPARATION COMPOSITE YIELDS	SP.GR.	.9165		. 9165		DETAILED MATERIAL BALANCE L.P. HYDROTREATING COMPOSITE YIELDS	SP.GR.	9165	1.0000	1	1.0000	. 8673	1	DETAILED MATERIAL BALANCE H.P. HYDROTREATING COMPOSITE YIELDS	SP.GR.	. 6673	1.0000		1.0000	
:						:		,					:			1			:	
	# C # C # C # C # C # C # C # C # C # C	OCCIDENTAL SHALE OIL	CHARGE	PRODUCTS DEASHED SHALE OIL	PRODUCTS		,	DEASHED SHALE OIL	COLD TREATED WATER	CHARGE	PRODUCTS SOUR WATER - L.P.HT HYDDOGEN SUILETOE	SEPARATOR LIQUIDILP) LOSS	PRODUCTS			CHANGE SEPARATOR LIQUID(LP)	COLD TREATED HATER	CHARGE	PRODUCTS SOUR WATER - H.P.HT SEPARATOR LIQUID(HP) LOSS	PRODUCTS

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMLM JP-8 JET FUEL

		MSCFO	0000	26.50			29761.69			i		:	HSCFO		•	AC 788		
		ž	93.49	3.24	100.00		4.22	19.97	4.54	90.	100.00	i	אַנא	100.00	100.00	19.	68.72 29.65	100.00
ļ		TONS/D	13195.34	456.95	14113.74	+	596.03	2626.40	10363	8.58	14113.74	:	TONS/0	2626.40	2626.40	18 67	1804.78	26.26 40 100 00
MCE		Ľ	97.18	2.82	100.00	1	à	\$ 0 · \$ 7	3.95	72.487	107.28	ANCE	ראג	100.00	100.00		72.34	97.73
'ERIAL BALANCE IG	ELDS	887	89966.8	2609.9	92576.7	!		1.54422	3659.0	7	99320.0	FERIAL BAL ITTER IELDS	8870	22993.1	22993.1		16633.9	22671.4
DETAILED MATERIA	COMPOSITE YIELD	SP. GR.	7118.	1.0000				\$260·	1.0000			DETAILED MATERIAL BALANCE NAPHTHA SPLITTER COMPOSITE YIELDS	SP.GR.	.6524	•		.6197	
	•	u e e e e e e e e e e e e e e e e e e e	SEPARATOR LIQUID(HP)	COLD TREATED WATER EDE STRIPPING STEAM	CHARGE	PRODUCTS	LP FLASH GAS: JP-8	PRUD PRACT DV: JP-8	SOUR MATER (MC)	LOSS	PRODUCTS			CHANGE PROD FRACT OV: JP-8	CHARGE	PRODUCTS SDITTED GAS ID-A	SPLITTER BOTTOMS	STUIDOGG

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

	DEBLITANT YED	בעלאר משה					
	COMPOSITE YIELDS	ELDS		:			
CHABGE	SP.GR.	88170	Ľ	TONS/D	MT.	HSCFD	
SPLIT OV LIG=JP-8	.6197	16633.9	100.00	1804.78	100.00		
CHARGE		16633.9	100.00	1804.78	100.00		
PRODUCTS DEBUT OVER'D=JP-8 LT NAPHTHA JP-8	5458	6263.9	37.66 62.35	598.59 1206.19	33.17	8664,78	
PRODUCTS		16634.5	100.00	1804.78	100.00		
	DETAILED MATERIAL BA DEPHOPANIZER COMPOSITE YIELDS	20	LANCE	:	i		Apper
A DOCA	SP.GR.	B8/0	ĽĶ	TONS/D	X X	HSCFD	ndix
DEBUT OVER'D=JP-8	.5458	1229.3	100.00	117.47	100.00	1739.69	B .
CHARGE		1229. 3	100.00	117.47	100.00		.7 (
PRODUCTS DEPROP OV'D JP-8 MIXED C4'S:JP-8	.5743	9.078	68.38	32.95 84.52	28.05 71.95	637.45	(Cont.)
PRODUCTS		940.6	68.38	117.47	100.00	and the second s	

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

	DETAILED MATERIAL BALANCE FUEL GAS TREATER	TERIAL BAL	ANCE				
	COMPOSITE YIELDS	IELDS	:	 - 			
CHARGE	SP. GR.	8870	ראג	TONS/D	77	MSCFD	
LP FLASH GAS: JP-8 DEBUT OVER: 0=JP-8 SPLITTE GAS=JP-8	8575.	5034.7	100.00	596.03 481.12 42.81	50.67 40.90 3.64	29761.69 7125.09 887.76	
NAP HYDROTREAT GAS F.O. STAB OVER'D					2.80 .06 1.92	617.45 102.21 1235.32	
CHARGE		5034.7	5034.7 100.00	1176.39	100.00		
PRODUCTS HYDROGEN SULFIDE TRI 1 P FIASH: 10-8				18.63	1.58	414.81	Арр
TRI DEB OV'D=JP-B				476.74	47. /8	7027.35	
				41.90	3.56 2.56	667.76	di
18 TRI NAP HTRT GAS	1			86.	90	102.21	K B
				20.94	1.78	1199.58	·• /
PRODUCTS				1176.39	100.00		((
	DETAILED MATERIAL BALANCE SULFUR PLANT COMPOSITE YIELDS	TERIAL BAL T IELDS	ANCE	:	1		Cont.)
	SP. GR.	88170	ľ.	TONS/D	**	MSCFD	
HYDROGEN SULFIDE				96.20	100.00	2186.99	
CHARGE	!			98.20	100.00		
PRODUCTS SULFUR SULFUR PLANT LOSS				87.77 10.43	89.38 10.62		
PRODUCTS				98.20	100.00		

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL

		A WAR AND A TOTAL AND ADDRESS OF THE PARTY O								A C C. Company Company and make paying Company depth common to the Company of the				Apper	ad i x	: В	•7 (Cor						
	MSCFD	F11 24	20520 27	7027.15	AC 7.48	613.61	10 201	1100 FA	0000 E	285.78				234287.62				MSCFD	1	169.79			102.21	
	HTZ.	8	70.7	36.56	40	2.31					9.04	100.00	ı	54.50	100.00		•	HT.	8.14	3.79	100.00		3.79 3.79 %.10	100.00
	TONS/D	97.00	585.36	476.74	00.19	31.88	6	20.96	71 17	11.66	110.%	1380.43		752.27 628.16	1380.43		:	TONS/0	778.81	30.68	810.04		.93 30.68 778.42	810.04
ANCE	LVX	21 27	;								76.73	100.00				ANCE		Ľ	97.09	2.91	100.00		2.91	100.00
L BAL	881/0	289.2	:								954.0	1243.3				ERIAL BALA	OTREATER ELDS	681/0	\$837.5	175.3	6012.7		175.3	6012.9
DETAILED MATERIAL BAL HYDROGEN PLANT COMPOSITE YIELDS	SP. GR.	9415				1					.6643					DETAILED MATERIAL BAL	NAPHTHA HYDROTREATER COMPOSITE YIELDS	SP. GA.	. 7620	1,0000 175.3			1.0000	
	CHARGE	DEBUT OVERHEAD-100R	TRT LP FLASH: JP-8	TRT DEB OV'D=JP-8	TRT SPLIT GAS=JP-8	TRT DEP OV'D-JP8	TRT NAP HTRT GAS	TRT F.O. STAB OV'D	PLAT NET SEP GAS-100	PLAT DEBUT GAS-100R	LT NAPHTHA JP-8	CHARGE		PRODUCTS HYDROGEN (972) CO2HYD.PLT.	HODUCTS 599	-			SPLITTER BOTTOMS	COLD TREATER	CHARGE	PRODUCTS	NAP HYDROTREAT GAS SOUR WATER-NAP HT TRT HC HVY NAPHTHA	PRODUCTS

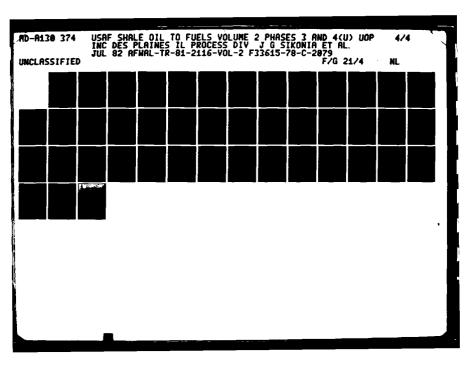
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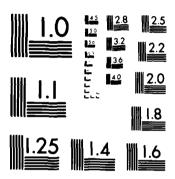
UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 3 - MAXIMLM JP-8 JET FUEL

			The second secon			Ap	pendix	B.	1 (Con	t.)							The state of the s	
		MSCFD			8928.54 285.78 531.26			MSCFD					:			HSCFD	2186.99		2186.99
		ž	100.00	100.00	9.40 1.47 3.48 85.65	100.00	!	¥	19.85	20.41	100.00	90.54	. 18	100.00		MTX.	100.00	100.00	100.00
	· •	TONS/0	778.42	778.42	73.17 11.44 27.09 666.71	778.42		TONS/D	623.02	640.63	3138.88	2841.97	5.57	3138.68		TONS/D	98.20	98.20	98.20
ij	•	۲۷	100.00	100.00	4.95	84.19	ij	ראג	19.65	20.41	100.00	90.54	· · · · · · · · · · · · · · · · · · ·	90.54	7	ראא			
RIAL BALANCE	son	8870	5837.6	5837.6	289.2	4914.5	RIAL BALAN EATING LDS	B8 C0	3558.4	3659.0	17927.7	16231.9		16231.9	RIAL BALAP ATION LDS	BB/0			
DETAILED MATERIA	COMPOSITE YIELDS	SP. GR.	. 7616		. 5349		DETAILED MATERIAL BALANCE SOUR MATER TREATING COMPOSITE YIELDS	SP. GR.	1.0000	1.0000		1.0000	1 1 2 3 4 4		DETAILED MATERIAL BALANCE AMINE REGENERATION COMPOSITE YIELDS	, p. GR.			1
		u 0 2 1	TRT HC HVY NAPHTHA	CHARGE	PRODUCTS PLAT NET SEP GAS-100 PLAT DEBUT GAS-100R DEBUT OVERHEAD-100R 100 RON PLATFORMATE	PRODUCTS		- AABGE	SOUR HATER - L.P.HT	HATER		PRODUCTS TREATED SOUR WATER APPROATA	\$50	PRODUCTS		1	301a	# · ·	w 33 34 34

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIHIZATION REPORTS, P51.0

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USAF SHALE OIL TO FUELS Phase IV Case 3 - Paximim JP-8 Jet Fuel

COMPOSITE YIELDS COMPOSITE YIELDS PRODUCTS TREATED SOLE HATER		DETAILED MATERIAL APPONIA PLANT		BALANCE				1 1 1
SP.GR. BBLD LVZ TONS/D HTZ CHARGE CHARGE CHARGE 1.0000 43.6 291.34 100.00 BETAILED HATERIAL BALANCE FUEL OIL STABILIZED CHARGE		COMPOSITE Y	recos		•	,		
STATE STAT	HARGE	SP. GR.	887	Ľ	TONS/D	F 4	MSCFD	
STATE SOLF HATER 1.0000 83.8 14.66 5.03	APPONIA				291.34			i i
STATED SOLR HATER 1.0000 43.6 14.66 5.03	CHARGE				291.34			
PRODUCTS B1.6 291.34 100.00	PODUCTS TREATED SOUR MATER ANHYDRUS AFFONIA	1.0000	83.8		14.66	5.03		
SPECIFIC PATERIAL BALANCE COMPOSITE YIELDS SP.GR. BBL/D LVZ TGNS/D HTZ SP.GR. BBL/D LVZ TGNS/D HTZ SP.GR. BBL/D LVZ TGNS/D HTZ LO. STAB GVER'D G458.7 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 1240.63 100.00 12441.92 100.00 12441.92 100.00 12441.92 100.00 12441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 100.00 124441.92 124.00 124.00 124441.92 124.00 124.00 124441.92 124.00 124.00 124441.92 124.00 124.00 124.0	PRODUCTS	:	83.8		291.34	100.00		
SP.GR. BBL/D LV. TONS/D HTX SPARATOR LIQUID(HP) 6377 8458.7 100.00 1240.63 100.00 CHARGE 6458.7 100.00 1240.63 100.00 CHARGE 6640 6052.2 95.19 1218.09 99.18 FRODUCTS 6640 6052.2 95.19 1218.09 99.18 FRODUCTS 6640 6052.2 95.19 1240.63 100.00 DETAILED HATERIAL BALANCE COMPOSITE YIELDS SP.GR. BBL/D LVX TONS/D HTX SP.GR. BBL/D LVX TONS/D HTX SP.GR. BBL/D 100.00 14441.92 100.00 CHARGE 9165 90000.0 100.00 14441.92 100.00 PRODUCTS 9165 90000.0 100.00 14441.92 100.00 PRODUCTS 9165 90000.0 100.00 14441.92 100.00		DETAILED HA	FERIAL BAL	ANCE	 - - - -	i :		Ap
SP.GR. BBL/D LVX TONS/D HTX CHARGE CHARGE CHARGE 1.0. STAB OVER'D CONSTAB BOTTOMS DETAILED MATERIAL BALANCE COMPOSITE YIELDS CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE SP.GR. BBL/D LVX TONS/D HTX CCIDENTAL SHALE OIL 1.0. 0.000 CHARGE CHARGE SP.GR. BBL/D LVX TONS/D HTX SP.GR. BBL/D 100.00 14441.92 100.00 CHARGE SP.GR. BBL/D 100.00 14441.92 100.00 CHARGE SPROUCTS SP.GR. BBL/D 14441.92 100.00 CHARGE SPROUCTS SPROUC		COMPOSITE Y	rems					pet
CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE O. STAB OVER'D O. STAB OVER'D O. STAB BOTTOMS BOS2.2 95.19 1240.63 100.00 CCHBINED MATERIAL BALANCE CCHBINED FACILITIES CCH	ARGE	SP.GR.	88/70	Š	TONS/D	¥	HSCFD	nd1:
CHARGE D. STAB OVER'D O. STA	SEPARATOR LIQUID(HP)	.6377	8458.7	100.00	1240.63	100.00		ж В
D. STAB OVER'D D. STAB BOTTOMS OETAILED MATERIAL BALANCE COMPOSITE YIELDS CHARGE	CHARGE		8458.7	100.00	1240.63	100.00		.7
PRODUCTS DETAILED MATERIAL BALANCE COPESITE YIELDS COPPOSITE YIELDS COPPOSITE YIELDS SP.GR. BBL/D LVZ TONS/D MTZ SP.GR. BBL/D LVZ TONS/D MTZ CHARGE CHARGE TOONOO 100.00 14441.92 100.00 FRODUCTS PRODUCTS DETAILED MATERIAL BALANCE COPPOSITE YIELDS SP.GR. BBL/D LVZ TONS/D MTZ SP.GR. BBL/D LVZ T		0,98.	8052.2	95.19	22.54		1235.32	(Cont.
DETAILED MATERIAL BALANCE COPENINED FACILITIES COPPOSITE YIELDS SP.GR. BBL/D LVZ TONS/D HTZ SP.GR. BBL/D LVZ TONS/D HTZ CHARGE CHARGE SIDENTAL SHALE DIL .9165 90000.0 100.00 14441.92 100.00 PRODUCTS	PRODUCTS		8052.2	95.19	1240.63	100.00)
SP.GR. BBL/D LVZ TONS/D HTZ CIDENTAL SHALE DIL CHARGE CHARGE CHARGE SP.GR. BBL/D LVZ TONS/D HTZ -9165 90000.0 100.00 14441.92 100.00 FRODUCTS 90000.0 100.00 14441.92 100.00		DETAILED MA COPPOSITE Y	TERIAL BAL SILITIES TELOS	ANCE	:	:		
CHARGE CHARGE	ų (SP.GR.	88170	Š	TONS/D	H	MSCFD	
CHARGE CIDENTAL SHALE OIL .9165 90000.0 100.00 14441.92 PRODUCTS 90000.0 100.00 14441.92	OCCIDENTAL SHALE OIL	. 9165	90000.0	100.00	14441.92			
CIDENTAL SHALE DIL .9165 90000.0 100.00 1441.92 PRODUCTS 90000.0 100.00 14441.92	CHARGE		900000	100.00	14441.92	100.00		
90000.0 100.00 14441.95	ODUCTS OCCIDENTAL SHALE OIL	. 916	900000	100.00	14441.92	100.00		
	PRODUCTS		90000	100.00	14441.92	100.00		!

778-42 100.00 778-42 100.00 73.17 9.40 8928.54 11.44 1.47 285.78 57.09 3.48 5.65 778.42 100.00 623.02 19.65 1844.54 56.76 640.63 20.41 30.68 100.00 2841.97 90.54 291.34 9.28 5.57 18 3138.88 100.00 70×S/D HTZ PSCFD 70×S/D HTZ PSCFD 98.20 100.00
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	:			MSCFD					
	•			17 %	100.00	100.00	100.00	100.00	
				TONS/0	84.52 100.00	84.52 100.00	84.52 100.00	94.52 100.00	
51.0	:	NCE NCE		K	100.00	100.00	100.00	840.6. 100.00.	
EPORTS, P		RIAL BALANCE	SOT	298	840.6	9.079	9.0.6	9.059	
STANDARD OPTINIZATION REPORTS, P51.0	ಪ	DETAILED MATERIAL DUTHY LAIT	COMPOSITE YIELDS	SP. GR.	.5743		.5743	:	
	FUELS JP-8 JET FU				; ; ; ;				ļ
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMUM JP-8 JET FUEL			CHARGE	MIXED BUTANES	CHARGE	PRODUCTS MIXED BUTANES	PRODUCTS	

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0

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	UNITS/0	9720.85	171627.72	117422.70	638483.09	8993.59	2327.09	147.51	į	2157.87	175873.57	4124.25	14233.36	6788.77	4003.95	938.71	1385.16		
	UNITATON	.673	11.884	9.160	48.387	3.424	1.289	1.256	1	21.974	233.790	5.2%	18.285	2.163	40.773	3.393	1.116		
104500/UNIT	UNIT/86L	.1080	1.9070	1.2677	7.0%9	. 3911	. 1399	.1200		•		. 7065	2,4382	. 3787			. 1638	•	
PRICE 15	TONS/0	14441.92	14441.92	14390.21	13195.34	2626.40	1604.78	117.47	1176.39	98.20	752.27	778.81	778.42	3138.88	36.20	276.68	1240.63	14441.92	84.52
F AN UNPOOLED UTILITY UNIT 15 KWH	887	90000.0	90000	92629.0	89966.8	22993.1	16633.9	1229.3		•		5837.5	5837.6	17927.7			8458.7	90000	9.00.6
DETAILED USAGE OF AN U. ELECTRIC POWER UNIT	CONTROL STREAM	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE
	MODE	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COPBINED HODES	COMBINED MODES
	PROCESS UNIT	FEED PREPARATION	L.P. HYDROTREATING	H.P. HYDROTREATING	HYDROCRACKING	NAPHTHA SPLITTER	DEBUTANI ZER	DEPROPANIZER	FUEL GAS TREATER	SULFUR PLANT	HYDROGEN PLANT	NAPHTHA HYDROTREATER	LOP PLATFORMING	SOUR WATER TREATING	AMINE REGENERATION	APPONIA PLANT	FUEL OIL STABILIZER	COMBINED FACILITIES	DUPIT UNIT

(Cont.)

PAGE 19			2.	and the state of t						1			•	A	P	en	d:	İx	:]	В.	7	2030.40	t •
			UNITS/D		1733.03	1462.05	576.92				21.57	-5999.61		-151.77							-457.81	4151.30 20	4141 01
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			UNIT/88L	!	.0193	.0158	5200							0603				!			j		
		} 1 ;	TONS/D	14441.92	14441.92	14390.21	13195.34	2626.40	1804.78	1176.39	96.20	752.27	778.81	778.42	3138.88	98.20	276.68	1240.63	14441.92	84.52	1		
1.0		OF A POOLED UTILITY LINIT IS HIBS	86170	90000	90000.0	92629.0	83966.8	22993.1	16655.9	7567.3			5637.5	5837.6	17927.7			8458.7	90000	9.058			
STANDARD OPTINIZATION REPORTS, PS1.0		ETAILED USAGE OOB STEAM OSTS ARE ALLOC	CONTROL STREAM	COMPOSITE USAGE	COMPOSITE USAGE				COMPOSITE USAGE			COMPOSITE USAGE		COMPOSITE USAGE	COMPOSITE USAGE		\$			COMPOSITE USAGE			
	USAF SHALE OIL TO FUELS PHASE IV CASE 3 - HAXTHM ID-A IFT FIEL		HOOE	COMBINED MODES				COMBINED MODES	CONSTRUCTIONS			COMBINED MODES	_	COMBINED MODES	COMBINED MODES					COMBINED MODES		EFINERY FUELS	A STRAM
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV CASE 3 - HAXIM H. ID-A II	1	PROCESS UNIT			H. P. HYDROTREATING	;	DEBLITANT SPLITTER	a	ATER	_	_	EATER	_		ATION	_	_	ACILITIES		TOTALS	ATOUNT PRODUCED FROM REFINERY FUELS	AMOUNT CONVEDTED TO JEAN STEAM

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AFOLNT CONVERTED TO 508 STEAM

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMIM JP-8 JET FUEL

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2		6 .30	-	107.36	20.60	5.77	3.16			18.18	2.07	14.85	71.65	95.42	8 0.8	5.09			1	12 81					
UNITS/D		145.86		3578.58	686.80	192.39	105.33			606.02	69.16	494.92	2395.10	3180.63	3032.67	169.59				16657.06					
UNIT/TON		010		.271	. 261	. 107	.897			909.	690.	419	. 763	32.389	10.81	137									
UNIT/BBL	•	. 0016		.039B	.0299	.0116	.0857				.0118	8480 ·	.1336			.0200									
TONS/D	14441.92	14441.92	14390.21	13195.34	2626.40	1604.78	117.47	1176.39	96.20	752.27	778.61	778.42	3138.69	98.20	276.68	1240.63	14441.92	84.52		!				•	
88/0	90000	90000	92629.0	B-9968	22993.1	16633.9	1229.3				5637.5	5837.6	17927.7			6458.7	90000.0	840.6							
CONTROL STREAM				COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE					COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITÉ USAGE	COMPOSITE USAGE		COMPOSITE USAGE	COMPOSITE USAGE							
HODE		_	_	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COPBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COPBINED MODES	COMBINED MODES	-							
PROCESS UNIT	FEED PREPARATION	L.P. HYDROTREATING	1. P. HYDROTREATING	HOROCRACKING	NAPHTHA SPLITTER	DEBUTANI ZER	DEPROPANI ZER	UEL GAS TREATER	WLFUR PLANT	DEPOSEN PLANT	APHTHA HYDROTREATER	OP PLATFORMING	OUR HATER TREATING	HINE REGENERATION	PHONIA PLANT	UEL OIL STABILIZER	Orbined FACILITIES	war wit							
	HODE CONTROL STREAM BBL/D TONS/D UNIT/BBL UNIT/TON UNITS/D	ESS UNIT MODE CONTROL STREAM BBL/D TONS/D UNIT/BBL UNIT/TON UNITS/D PREPARATION COMBINED MODES COMPOSITE USAGE 900000.0 14441.92	FSS UNIT MODE CONTROL STREAM BBL/D TONS/D UNIT/BBL UNIT/TON UNITS/D PREPARATION COMBINED MODES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66	ESS UNIT MODE CONTROL STREAM BBL/D TONS/D UNIT/TON UNIT/TON UNITS/D PREPARATION COMBINED MODES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66 4 HYDROTREATING COMBINED MODES COMPOSITE USAGE 926.29.0 14390.21 .0016 .010 145.66 4	COMBINED MODES	COMBINED MODES	COMPINED PROFES CONTROL STREAM BBL/D TONS/D UNIT/BBL UNIT/TON UNITS/D	COMPINED HODES	COMBINED HODES	COMPINED HODES	COMBINED HODES	COMBINED HODES	COMPINED HODES	COPEINED PROFES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66 4.36	COPENED PROFES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66 4.36 4.36 .0016 .0016 .0010 .002.34 .002.3	COPBINED PROPES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66 4.36 4.36 .0016 .010 145.66 4.36 4.36 .0016 .010	COPEINED HODES	COPEINED MODES COPPOSITE USAGE 900000.0 14441.92 .0016 .010 145.66 4.36	COPBINED PROCES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66	COPEINED HODES COMPOSITE USAGE 90000.0 14441.92 .0016 .010 145.66 4.38 .186	COPENED HODES	COPENIED HODES COMPOSITE USAGE 90000.0 14441.92 0.0116 145.66 4.38	COPENED HODES COMPOSITE USAGE 90000.0 14441.92 0.016 0.10 145.66 4.36	COPEINED PROSES COMPOSITE USAGE 90000.0 14441.92 0.016 0.10 145.66 4.36	COPENED FORES COPPOSITE USAGE 900000 14441.92 0016 010 145.66 4.38

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMLM JP-8 JET FUEL

		Appendix B.7	(Cont.)
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	UNITS/D	1961.21 4466.72 15551.15 1472.10 99.06 1890.11	42754.91 30417.39 3773.98
	UNITATION	.136 .1179 .1179 .560 .57.084 .127 .428	
	UNIT/BBL	. 0218 . 0462 . 1729 . 0640 . 0170 . 1218	
	10NS/0	14441.92 14490.21 14390.21 13195.34 2626.40 1804.78 117.47 117.47 1178.81 778.42 778.4	
S HETU S HETU WILITY POOL	0/198	90000.0 926.29.0 89966.8 22993.1 16633.9 1229.3 5837.5 5837.5 5837.5 6459.7	•
DETAILED USAGE OF A POOLED UTILITY REFINERY FUELS LAIT IS MEDIU COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM	COMPOSITE USAGE COMPOSITE USAGE	
; ; ;	HOOF	COTBINED MODES COTBINED MODES	PRODUCED FROM F.O. STAB BOTTONS PRODUCED FROM LT NAPHTHA JP-8 CONVERTED TO 600# STEAM
,	PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING NAPHTHA SPLITTER DEBUTANIZER DEPROPANIZER FUEL GAS TREATER SULFUR PLANT HYDROGEN PLANT HYDROGEN PLANT NAPHTHA HYDROTREATER UQP PLATORHING SOUR WATER TREATING AHTHE REGENERATION APTONIA PLANT FUEL OIL STABILIZER COPBINED FACILITIES OLITHY UNIT	AFOLNT PRODUCED FROM F.O. STAB AFOLNT PRODUCED FROM LT NAPHTH AFOLNT CONVERTED TO 6008 STEAM

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UNIVERSAL DIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMLM JP-8 JET FUEL

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	0/						
	UNITS/0			-745.65	4.05 4719.58 163.03	6161.01	4141.01
; ;	UNIT/TON			-7.593	1.504		
	UNIT/BBL			· •	. 2633		;
į	10NS/D	14441.92 14441.92 14390.21	2626.40 2626.40 1804.78 117.47	1176.39 98.20 752.27 778.81	778.42 3138.68 98.20 276.68	1441.92	
LED UTILITY S HLBS MILITY POOL	BBL/0	90000.0	89966.8 22993.1 16633.9 1229.3	5837.5	17927.7	90000.0	
DETAILED USAGE OF A POOLED UTILITY 150# STEAM UNIT IS MLBS COSTS ARE ALLOCATED TO UTILITY POOL	CONTROL STREAM	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE		COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE		
: !	MODE	COMBINED HODES COMBINED HODES COMBINED HODES			COPBINED MODES COPBINED MODES COPBINED MODES COPBINED MODES		600# STEAN
	PROCESS UNIT	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING	HYDROCRACKING NAPHTHA SPLITTER DEBUTANIZER DEPROPANIZER	FUEL GAS TREATER SULFUR PLANT HYDROGEN PLANT NAPHTHA HYDROTREATER	UOP PLATFORHING SOUR HATER TREATING AHINE REGENERATION APPONIA PLANT FUEL OIL STABLIZER	COPBINED FACILITIES DUPIT UNIT	AYOUNT PRODUCED FROM 600# STEAM

				a and the same of									Αį	рp	e1	nd	1:	ĸ	В	-7	' (Co	nt	: •)
			9/8																					
	:		UNITS/D				-278. 42		765.41	33,93		-968.63					915.88			;		66A. 17		468.17
			UNITATION				021		. 424	. 289		*9.864	1				9.327							
			UNIT/BBL				-, 0033		0950.	.0276					1									
		:	TONS/D	14461.92	14441.02	14390.21	13155.14	2626.40	1804.78	117.47	1176.39	98.50	752.27	778.61	778.42	3138.88	98.20	276.68	1240.63	14441.92	84.52	;		
	•	OF A POOLED UTILITY UNIT IS HIBS CATED TO UTILITY POOL	88	0.00006	90000.0	92629.0	89%67	22993.1	16633.9	1229.3	;			5837.5	5837.6	17927.7			6458.7	90000.0	9.058			
		DETAILED USAGE OF A PO SO# STEAM UNIT COSTS ARE ALLOCATED TO	CONTROL STREAM	COMPOSITE USAGE			COMPOSITE USAGE	COMPOSITE USAGE						COMPOSITE USAGE	COMPOSITE USAGE				COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE			
USAF SHALE OIL TO FUELS PHASE IV	CASE 3 - MAXIMUM JP-8 JET FUEL	<u> </u>	HODE	COMBINED MODES	COPBINED MODES		COMBINED MODES												COPBINED MODES	COMBINED MODES	COMBINED MODES			600# STEAH
USAF SHALE	CASE 3 - MA		PROCESS UNIT	FEED PREPARATION	L.P. HYDROTREATING	H. P. HYDROTREATING	HYDROCHACKING	NAPHTHA SPLITTER	DEBUTANIZER	DE PROPANI ZER	FUEL GAS TREATER	SULPUR PLANT	HIDROGEN PLANT	NAPHTHA HYDROTREATER	UN PLAIFORMING	SOUR MATER TREATING	AMINE REGENERATION	AFFICINIA PLANT	FUEL OIL STABILIZER	COMBINED FACILITIES	DUMY UNIT	TOTALS	1	AMOUNT PRODUCED FROM 600# STEAM
			PROCES	FEED PA	¥ .d .i	I Q	H) DROC	MANTH	OEBUTA	DEPROP	֓֞֝֝֞֝֞֝֝֞֝֝֓֞֝֞֝֓֓֞֝֞֝֓֓֓֞֝֓֓֓֓֞֝֓֓֓֞֝֓֓֞֝֓֓֞֝֓֓֞֝֓֓֞֝֓֡֓֞֝֓֡֓֓֡֝֓֡֝֓֡֝֞֝֓֡֓֞֝֡֓֡֓֝֡֝֓֡֝֝֡֓֡֝֝֡֡֝֝֡	301708	HOROE	MAPHTH	3	anos	ATINE		i	7	_	TOTALS		

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMIM JP-8 JET FUEL

UNIVERSAL OIL PRODUCTS CO.

		Appendi	ж В	•7	(Cont.)
	0/•			2780.43	
	UNITS/D	1006.80 1860.52 12854.46 576.18	16297.97	5560.86	10737.11
	UNIT/TON	18.946 17.088			•
	UNIT/88L	.0987	!		· :
,	10NS/0	14441.92 14490.21 13195.34 26.56.40 1804.78 1176.47 1176.47 778.81 778.81 778.81 778.82 778.81 778.68 118.68 98.20 752.27 778.81 778.42 84.52 14441.92			
POOLED UTILITY IT IS MLBS TO UTILITY POOL	8870	90000.0 92629.0 92629.0 89966.8 22993.1 16633.9 1229.3 5837.8 5837.8 5837.8 17927.7			
DETAILED USAGE OF A BOILER WATER UNIT	CONTROL STREAM	COMPOSITE USAGE COMPOSITE USAGE			
	HODE	CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES CO-BINED HODES			CONDENSATE
i	PROCESS UNIT	L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING HYDROCRACKING HYDROCRACKING HYDROCRACKING HYDROCRACKING DEBUTANIZER DEBUTANIZER FUEL GAS TREATER SULFUR PLANT HYDROGEN PLANT HYDROGEN PLANT NAPHTHA HYDROTREATER UOP PLATFORHING SOUR WATER TREATING ANTINE REGENERATION ATTONIA PLANT FUEL OIL STABILIZER COPBINED FACILITIES DUMNY UNIT	TOTALS	AMOUNT PURCHASED	AYOUNT FRODUCED FROM CONDENSATE

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0

USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMAN JP-8 JET FUEL DETAILED USAGE OF A POOLED UTILITY CONDENSATE UNIT IS HLBS

			COSTS ARE ALLOCATED TO UTILITY POOL	UTILITY POOL						
	PROCESS UNIT	MODE	CONTROL STREAM	88170	TONS/D	UNIT/BBL	UNIT/TON	UNITS/D	0/4	
	FEED PREPARATION	COPBINED MODES	COMPOSITE USAGE	90000	14441.92	•		1	eri ener i descripti descripti de la como de	:
	L. P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	90000.	14441.92	0193	120	-1733.03		
	H.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.	14390.21	0158	-, 102	-1462.05		
1	HIDROCRACKING	COMBINED HODES		89966	13195.34	- 0075	150.1	-676.92		
	NAPHTHA SPLITTER	COMBINED MODES		22993.1	2626.40	•				
	DEBUT ANI ZER	COMBINED MODES	COMPOSITE USAGE	16633.9	1804.78	0460	424	-765.41		
	DEPROPANI ZER	COMBINED MODES	COMPOSITE USAGE	1229.3	117.47	0276	289	-33.93		
	FUEL GAS TREATER	COMBINED MODES	COMPOSITE USAGE		1176.39				***************************************	!
	SULFUR PLANT	_			98.20		684	-67.13		
	HIDROGEN PLANT	COMBINED MODES			752.27					
	MAPHTHA HYDROTREATER	COMBINED HODES	COMPOSITE USAGE	5837.5	778.81					
	LIOP PLATFORMING		COMPOSITE USAGE	5837.6	778.42	- D343	-,257	-200.11		
	SOUR HATER TREATING		COMPOSITE USAGE	17927.7	3138.88	2633	-1.504	-4719.58		ΑĮ
	AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.20		-10.987	-1078.94	•	р
-2	_	COMBINED MODES	COMPOSITE USAGE		276.68					er
27		COMBINED MODES	COMPOSITE USAGE	8458,7	1240.63				- 140 C C C C C C C C C C C C C C C C C C C	nd
9-	_	COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92					1>
-	DUMMY UNIT	COMBINED MODES	COMPOSITE USAGE	9.078	84.55					c F
i	TOTALS				,			-10737.11		3 . 7
	AMOUNT CONVERTED TO BOILER WATER	BOILER HATER						10737.11		(Cont
			1	!		1	· : : : :			•)

STANDARD OPTIMIZATION REPORTS, P51.0 UNIVERSAL DIL PRODUCTS CO.

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USAF SHALE OIL TO FUELS PHASE IV CASE 3 - MAXIMJM JP-8 JET FUEL

DETAILED USAGE OF AN UNPOOLED UTILITY
CAT. & CHEMICALS LAIT IS \$
COSTS ARE ALLOCATED TO PROCESS UNITS

Appendix B.7 (Cont.) 16.67 10308.64 3858.02 6290.22 3 27.75 3775.67 16.59 331.99 10308.64 3858.02 6290.22 27.75 UNITS/D 5.019 .021 .426 WITTIN 268 . 283 PRICE IS 1.00000/LINIT .1145 0028 UNIT/BBL 14441.92 14390.21 13195.34 2626.40 110.39 110.39 110.39 752.27 778.81 778.42 3138.88 20.20 276.68 12461.92 TONS/D 14441.92 90000.0 90000.0 92629.0 89966.8 22993.1 16633.9 5837.5 5837.6 17927.7 9458.7 90000.0 840.6 8612 COHPOSITE USAGE
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L. P. HYDROTREATING
H. P. HYDROTREATING
HYDROCRACKING
NAPHTHA SPLITTER
DEBUTANIZER
DEPROPANIZER
FUEL GAS TREATER
SULFUR PLANT UOP PLATFORHING SOUR WATER TREATING FUEL OIL STABILIZER COMBINED FACILITIES AMINE REGENERATION HIDROGEN PLANT APPONIA PLANT PROCESS UNIT CHY CNIT

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, PS1.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 3 - MAXIMLM JP-8 JET FUEL

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3	PROCESS UNIT UTILITY CONSUMPTION SUPPARY
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JET FUEL	Æ
S JET FUEL	æ
-8 JET FUEL	æ
P-8 JET FUEL	æ
JP-8 JET FUEL	*
n JP-8 JET FUEL	*
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ITUT JP-8 JET FUEL	8
XINUM JP-8 JET FUEL	8
MXIMUN JP-8 JET FUEL	8
MAXIMUN JP-8 JET FUEL	8
- reasings JP-8 JET FUEL	8
- raxina JP-8 JET FUEL	**
s - naxinun JP-8 JET FUEL	**
ise s - naxinun JP-8 JET FUEL	**

		Appendix B.7 (Cont.)
CONDENSA TE HLBS	-1733.0 -1462.0 -1626.9 -765.4 -33.9 -67.1 -47.9	-1078.9
BOILER M ATER MLBS	1006.8 1860.5 12854.5 576.2	16298.0
SO# STEA M MLBS	-278.4 765.4 33.9 -968.6	468.2
150# STE AH MLBS	-745.7 -75.7 4.079.8	163.0
REFINERY FUELS MMBTU	1961.2 9466.7 15553.4 1472.1 43544.3 99.1 1890.3	411.3
COOLING HATER MGAL	145.9 1578.6 686.8 192.4 105.3 606.0 69.2 494.9	3180.6 3032.7 169.6 14657.1
600# STE AM MLBS	1733.0 1462.0 676.9 21.6 -3999.6	-457.8
ELECTRIC POWER KWH	9720.9 171627.7 117422.7 638483.1 8993.6 2327.1 147.5 2157.9 17587.9 4124.2 14233.4 6788.8	4004.0 938.7 1385.2 1158228.2
PROCESS UNIT	L.P. HYDROTREATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING HYDROCRACKING HOPHINA SPLITTER DEBNOTANIZER DEFROPANIZER SULFUR PLANT HYDROGEN PLANT HYDROGEN PLANT NAPHTHA HYDROTREATER UOP PLATFORMING SOUR HATER TREATING	AHINE REGENERATION ATHONIA PLANT FUEL OIL STABILIZER TOTAL CONSUMPTION

CASE 3 - M	USAF SHALE DIL TO FUELS PHASE IV CASE 3 - MAXIMIM JP-8 JET FUEL		; ; ;		:	:
		PROCESS UNIT UTILITY CONSUMPTION SUPPLARY	TION SUPPLARY			
PROCESS UNIT	CAT. & C HEMICALS					
FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING	10308.6 3858.0 6290.2		: :			
naphtha splitter Debutanizer Depropanizer Sulfur Plant						
HYDROGEN PLANT NAPHTHA HYDROTREATER UOP PLATFORMING SOUR HATER TREATING	3775.9 16.6 332.0			:		, A
AHINE REGENERATION APPONIA PLANT FUEL OIL STABILIZER	27.8					opendi:
TOTAL CONSUMPTION	24609.1			•		
; ; ; ; ,						

JP-8 PLUS DIESEL CASE -- STANDARD OPTIMIZATION REPORT

CALVERSA CALVERSA	UNIVERSAL DIL PRODUCTS CO.	STANDARD OFTIMIZATION REPORTS, PS1.0	PORTS, PS	3.0			15:11 NOV 20. A1	A) 0ACF	•
	USAF SHALE OIL TO FUELS PHASE IV								
	CASE 4 - JP-8 JET	CASE 4 - JP-6 JET FUEL + OF-2/OFH DIESEL	i			:			
		GROSS HARGIN SLITMRY	LIFTHEN						
SALES	MARKET	PRODUCT	HADDAL	QUANTITY LIMITS	FIXED	QUANTITY SOLD	PRICE	TOTAL	1
	44	UNERDED GASOLINE				\$169.30 BBL	58.8500	186	
		AV IUMBINE PUEL JP-B OF-2/OF-M DIESEL				4%697.16 BBL 26894.99 BBL	57.85.00	2914679.04	i
-	. ⋖	ANYDRIS ATENCA			;		95.4500	6176.17	
	TOTAL SALES REVENUE							474500.78	İ
RAH HATE	RAH MATERIAL PURCHASED	MATERIAL		QUANTITY LIMITS	9	GUNNTITY PURCHASED.	PRICE	TOTAL	ì
	•	OCCIDENTAL SHALE DIL COLD TREATED HATER -508 STRIPPING STEAM		.	0.0000	90000.00 BBL 14504.43 BBL 159.99 ION	40.0000 .0245	3600000.00	
•	TOTAL RAH MATERIAL COST	COST						TABATER 14	
GROSS MAGIN	AGIN		!	:				1144744.92	

USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

UNIVERSAL OIL PRODUCTS CO.

OPERATING COST SUPPARY

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100				38 26473.23	20 22134.96			į			_		è	77 1373.20	50 3065.68	53 1649.15	1749.01		_			157848.21			4912.06			-•	:			514.55	
ODEDATING COST			:		4 1.53620	Ī				-	06726	19.67463	Ā		0 8.88260	1.18263	17.81405	2.68881		•	!					•			2.	•	514.65		
!	\$/BB	1000	024220	. 294147	. 238964	.627371	08195¢	110762	37000	. 0000			1	. 530560	1.184450	. 207060			77477	09805			1		;			•	•				
CHIANTITY	TOVD		26.14441	14441.92	14390.21	12450.46	1338.05	911.18	0	77.7	60.34	40. 10 40. 10	715.82	145.31	345.13	3085.64	98.18	275.88	1985.51	14441.92	40.78		?	4912.06		9,4	8		SOURCE	}	REFINERY FUELS		
į	85	0000	40000	90000	92629.0	64888.1	11848.5	86.76.8	415					2288.2	2588.3	17623.6			13537.4	90000.0	405.4		#ZUNIT	.5000		*/UNIT	. 00010						
							į									1 1							UNITS/D	9824.12		UNITS/D	625.78		0/STIND		754.67		
CONTROL STREAM					SITE COST	SITE COST	SITE COST								_	DMPOSITE COST.	DMPOSITE COST	SITE COST	SITE COST	SITE COST					:				LNIT		MLBS	:	
	}	Š	3170255	5	2	200	COMPC	COMPO			֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓	֡֝֞֝֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֡֓֓֓֓֓֓֡֓֓֡	COMPOSITE	5	0 7	COMPO) PEO:	S P P P P P	CHOU	COMPO	COMPOSITE		באַז	MLBS		13	MLBS		2		Ξ.	!	
, W	•	CHRINED MODES				COMBINED MODES	COMBINED MODES		COMBINED MODES		_		CONDINED HOUSE				COMBINED MODES	COMBINED MODES	COMBINED MODES	COPBINED MODES	COMBINED MODES	G COST	ITY	BOILER MATER		ITY	50# STEAM		UILITY	!	600# STEAM	srs	
HOOF		č	3 8	5	Š	S	9	5	Š	ב ב	į	3 8		_	_		5	_		_	8	OPERATIN	UTILITY	BOIL	ASES	WILITY	50%		COSTS			CTION CS	
PROCESS UNIT		FEED PREPARATION		THE PROPERTY OF THE PARTY OF TH	H. P. HICHOINEALING	HIDROCRACKING	NAPHTHA SPLITTER	DEBUTANI ZER	DEPROPANI ZER	FUEL GAS TREATER	SULFUR PLANT	HYDDOCEN DI ANT	NABHTHA MYDBOTOSATER	TOTAL TIONOLINE	OF PLAIFORNING	SOUR MAIER INEALING	AMINE REGENERATION	AMMONIA PLANT	FUEL OIL STABILIZER	COMBINED FACILITIES	DUM UNIT	TOTAL PROCESS UNIT OPERATING COST	UTILITY PURCHASES		TOTAL UTILITY PURCHASES	UTILITY SALES		TOTAL UTILITY SALES	UTILITY PRODUCTION COSTS			TOTAL UTILITY PRODUCTION COSTS	

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UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0 USAF SHALE OIL TO FUELS

USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL CAPITAL COST SUPPLARY

PROCESS UNIT	HOOF	CONTROL STREAM	3	ANTITY		APITAL CO	ST	
			BBC0 TON	101/10	\$/BBL	#/TON	?	
FEED PREPARATION	-		90000.0	14441.92		.59%0	8659.38	
L.P. HYDROTREATING	_		90000	14441.92		3.94621	56990.78	
H.P. HYDROTREATING	-		92629.0	14390.21		6.42342	92434.31	
HYDROCRACKING	_		64868.1	12450.46		13.01071	161989.29	
NAPHTHA SPLITTER	_		11848.5	1338.05	ł	1.15560	1546.25	- ;
DEBUTANI ZER	•		8676.8	933.38	. 314447	2.92312	2728.40	
DE PROPANIZER	-		632.9	60.19		10.93832	659.33	
FUEL GAS TREATER	•			766.39		1.56139	1196.64	
SULFUR PLANT	_			98.18	•	57, 38291	5633,94	
HYDROGEN PLANT	_			715.82		163.08393	116738.64	
NAPHTHA HYDROTREATER	-		2588.2	345.31	.775609	5.81350	2007.44	
UOP PLATFORMING	-		2588.3	345.13		17.18633	5931.57	
SOUR HATER TREATING	_		17623.6	3085.64	- :	3.02621	9337.79	
AMINE REGENERATION	COMBINED MODES	COMPOSITE COST		98.18		26.64689	2616.23	
APPONIA PLANT	-			275.88		1.50753	415.90	
FUEL OIL STABILIZER	_		13537.4	1985.51		1.46048	2899.80	
COMBINED FACILITIES	-		90000.0	14441,92	.115235	.71813	10371.11	
DUTHY UNIT	COMBINED MODES		405.4	40.78				

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REPORTS
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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

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CHARGE	SP.GR.	88	רא	TONS/D	25		
OCCIDENTAL SHALE OIL COLD TPEATED WATER 50# STRIPPING STEAH	.91650	90000.0	8 %	3, 2	84.252 14.815		
TOTAL CHARGE		104504.4	100.000	17141.42	100.000		
PRODUCTS	SP. GB.	: 298	ž į	TONS/D	X	ULDM	
UNLEADED GASOLINE AV TURBINE FUEL JP-8 DF-2/DF-M DIESEL SULFUR AWHIDRUS AFFONIA	,76091 ,60500 ,60500	3169.3 49697.2 26895.0	3.033 47.555 25.736	422.23 7004.50 3935.71 87.75	2.463 40.863 22.960 .512		
TOTAL PRODUCTS SOLD		79761.5	76.324	11726.08	1.609		
STREAMS CONVERTED TO UTILITIES	8 .6	5	Š	10NS/D	4.	0.054	<u> </u>
F.O. STAB BOTTONS	.86400	12886.8	12.331	1949.44	11.373		
TOTAL STREAMS CONVERTED		12886.8	12.331	1949.44	11.373		
STREAMS NOT UTILIZED	8.8	28	ř.	TONS/O	HTX		
LOSS SULFUR PLANT LOSS CO2HYD.PLT.				27.04	. 156	-	
	1.00000	16016.2	15.326	2804.20	3.642 16.359		
TOTAL NOT UTILIZED		16016.2	15.326	3465.90	20.219		
TOTAL PRODUCTS HADE		108664.5	103.961	17141.42	100.000		

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SMALE OIL TO FUELS
PHASE IV
CASE 4 - JP-8 JET FUEL + 0F-2/DFM DIESEL

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TIND	CAPACITY CATEGORY	CATEGORY	MAXIMUM	HINIH	FIXED	LP INPUT	ACTUAL	INVESTMENT	: ! !
FEED PREPARATION	CAPACITY	BBLS/CD				90000.0	90006	9030000	
L.P. HYDROTREATING	CAPACITY	BBLS/CD				900006	6.40006	59430000	
H.P. HTDROTREATING	CAPACITY	BBLS/CD		•		92630.0	92635.6	96390000	
HYDROCRACKING	CAPACITY	BBLS/CD				84887.5	84887.5	168931683	
NAPHTHA SPLITTER	CAPACITY	BBLS/CD				11849.6	11849.6	1612516	
DEBUT ANI ZER	CAPACITY	BBLS/CD				8677.4	8677.6	2845128	
DE PROPANI ZER	CAPACITY	BBLS/CD				636.0	636.0	686548	
FUEL GAS TREATER	CAPACITY	TONS/CD				766.4	766.4	1247922	
SULFUR PLANT	CAPACITY	TONS/CD				88.2	98.2	5875389	
HIOROGEN PLANT	CAPACITY	TONS/CD				715.8	715.8	121741719	i
NAPHTHA HYDROTREATER	CAPACITY	BBLS/CD				2588.2	2568.2	2093469	
UOP PLATFORMING	CAPACITY	BBLS/CD				2479.6	2479.6	6185784	
SOUR MATER TREATING	CAPACITY	BBLS/CD				17714.1	17714.1	9737981	
AMINE REGENERATION	CAPACITY	TONS/CO		i		28.5	98.2	2728353	1
APPONIA PLANT	CAPACITY	TONS/CD				275.9	275.9	433723	
FUEL OIL STABILIZER	CAPACITY	BBLS/CD				13537.2	13537.2	3024072	
COMBINED FACILITIES	CAPACITY	BBLS/CO				90000	90004	10815000	

UNIVERSAL OIL PRODUCTS CO. STANDARO OPTIMIZATION REPORTS, PS1.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

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UNIVERSAL DIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0	USAF SHALE OIL TO FUELS PHASE IV	CASE 4 - JD-A JET BIE: 4 PE-44555 STATES
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PRODUCT BLENDING

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AV TURBINE FUEL JP-8	8870		SPGR260F	MTX SULF	TONS/D SPGRAGOF WIX SULF VLX AROM SMOKE PT FREEZ PT VLX NAPH 7LASH PT	SMOKE PT	FREEZ PT	VC. NAPH	FLAST PT	D+L 401F	
HC KEROSINE: 300-550F	4%97.2	7004.50	.8050	.0003	10.0000	24.0000	/0.0000 24.0000 -58.0000	1.5000	100.0000	40.0000	
BLEND	49697.2	7004.50	.6050	. 0003		24.0000	-58.0000	:	1.5000 100.0000		
SPECIFICATIONS	•	HEN	.6398	. 4000	.4000 25.0000	20.0000	-59.0000	3.0000	3.0000	10.000	
	ř	PRODUCT BLENDING	DING								
AV TURBINE FUEL JP-8	88170		TONS/0 0+L 572F				l	,			
HC KEROSINE: 300-550F	49697.2	7004.50	100.000				I	•		•	
BLEND	4%97.2	7004.50	49692.2 7004.50 100.0000				;		:		
SPECIFICATIONS		AN	100.0000								
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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

	æ	PRODUCT BLENDING	DING							
DF-2/OF-M DIESEL	: B8/70	TONS/0	TONS/D SPGRa60F	MTX SULF	HT SULF FLASH PT CETANE I D+1 700F VIS 100F POUR PT.	CETANE I	D+L 700F	VIS 100F	POUR PT.	CLOUD PT
MC DIESEL:550-700F	26895.0	1935.71	. 8358	. 0003	286.0000	26.0000	100.0000	100.0000 18.4000	. 0000	5.0000
BLEND	26895.0	1935.71	.6358	. 0003	286.0000	56.0000	56.0000 100.0000 18.4000	16.4000	.0000	5.0000
SPECIFICATIONS	,	AT	.8607	. 7000	133.0000	45.0000	100.0000	100.0000 18.4100	0000	5.0000
	æ	PRODUCT BLENDING	DING							
DF-2/DF-M DIESEL	887	TONS/D	TONS/D D+L 675F	0+L 725F				-	:	1
HC DIESEL:550-700F	26895.0	3935.71	90.000	100.0000						
BLEND	26895.0	1915.71	3935.7190.000	100.0000		ı				
SPECIFICATIONS		AZ NIN	90.000	100.0000						

		•					
ENDING	D SPGRa60F	60	:		RECYCLE STREAM BLENDING	TONS/D SPGRa60F	5 .5744
PRODUCT BLENDING	BBL/0 TONS/D SPGR260F	275.98	275.88		RECYCLE ST	68L/0 TONS/	343.6 34.55 61.8 6.23
	ANHYDRUS AFFONIA	ANHYDRUS AFFICINIA	BLEND	NO SPECIFICATIONS		MIXED BUTANES	MIXED C4'S:JP-8 DH MIXED C4'S-98R PLT

405.4

NO SPECIFICATIONS

BLEND

TONS/D SPGR360F

87.75

NO SPECIFICATIONS

BLEND

PRODUCT BLENDING

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STANDARD OPTIMIZATION REPORTS, P51.0			BLEND FOR UTILITY PRODUCTION	TONS/D SPGRa60F	0,98.	. 8640	
IMIZATION		FM DIESEL	END FOR UT	TONS/D	1949.44	1949.44	
STANDARD OPT	UELS	UEL + DF-2/0	ฮ์	88/0	12886.8	12886.8	
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS PHASE IV	CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL		REFINERY FUELS	F.O. STAB BOTTOMS	BLEND	

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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/OFH DIESEL DETAILED MATERIAL BALANCE FEED PREPARATION COMPOSITE YIELDS

MTX MSCFD	100.00	100.00			100.00		HTX MSCFD	99.66	1.21 56852.12 3.13	100.00		.53 1772.18 95.32 .03	00.001		WTX HSCFD	38.34	1.70 86362.21 9.96	100.00	11.32	
TONS/D	14441.92	14441.92				}	10NS/D		182.55 472.83	15097.29		79.57 14390.21 4.48			TONS/D	14390.21		16289.43 10	1844.54 1 14435.97 8	4
רא	100.00	100.00	1 001	90 001			Ľ	97.09	2.91	100.00	3.64	99.92	103.76	ñ	ŗķ.	16.06	9.09	100.00	10.34	
88/0	90000	90000	0 00000	0.00000	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	ERIAL BALA EATING ELDS	88170	90000	\$700.6	92700.6	3558.4	92629.0	36187.4 101.76	D MATERIAL BALANCE DROTREATING TE YIELDS	8670	92629.0	9263.6	101892.6	10535,1 98425.5	
SP.GR.	. 9165		5916		DETATIED MATI	DETAILED MATERIAL BALANCE L.P. HYDROTREATING COMPOSITE YIELDS	SP.GR.	9165	1.0000		1.0000	.9673		DETAILED MATERIAL H.P. HYDROTREATING COMPOSITE YIELDS	SP. GR.	.6673	1.0000		1.0000	
CHARGE	OCCIDENTAL SHALE DIL	CHARGE	PRODUCTS DEASHED SHALE OIL	PRODUCTS			CHARGE	DEASHED SHALE OIL HYDROGEN (972)		CHARGE	PRODUCTS SOUR MATER - L.P.HT HYDROGEN SULFIDE	SEPARATOR LIQUID(LP)	PRODUCTS		CHARGE	SEPARATOR LIQUID(LP) HYDROGEN (972)	COLD TREATED WATER	CHARGE	PRODUCTS SOUR WATER - H.P.HT SEPARATOR LIQUID(HP) LOSS	

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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFH DIESEL

	DETAILED MATERIAL BALANCE HYDROCRACKING COMPOSITE VIELDS	TERIAL BALJ NG Telos			٠	1	
	3.45050	*****					
CHARGE	SP. GR.	88170	۲۸٪	TONS/D	MTX.	MSCFD	
SEPARATOR LIQUID(HP)	.8377	64888.1	97.18	, ~	93.63	! • • • • • • • • • • • • • • • • • • •	-
COLD TREATED WATER 50# STRIPPING STEAM	1.0000	2462.6	2.82	255.73 431.16 159.99	1.92 3.24 1.20	7%45.39	
CHARGE		67350.7	100.00	87350.7 100.00 13297.34	100.00		
PRODUCTS							
LP FLASH GAS: JPB-DM	•			406.51	3.06	25083.83	
HACO FRACT OV: JPB-DM	0549.		13.56	1338.05	10.06		
SOUR MATER (HC)	1.0000		3.95	604.47	4.55		
HC KEROSINE: 300-550F	6050	49697.2	56.89	7004.50	52.68		
MC DIESEL:550-700F	9318	0 10070	40				1

•	SP.GR. BBL/D SP.GR. BBL/D .6450 11848.5 100	BBC 11848.5	TER TDS DBC/O LV: 11848.5 100.00	TONS/D 1338.05 1338.05	LT.2. 100.00	HSCFD	
SPLITTER GAS=JPB-DH SPLIT OV LIQ=JPB-DH SPLITTER BOTTOMS	.6144	8676.8	73.23	25.80 933.38	1.93	601.35	
PRODUCTS		11516.6	97.20	1338.05 100.00	100.00		

PRODUCTS

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	MSCFD			4972.24	Andrew Andrews and the state of		HSCFD	754.48		303.93	79.68	
	MTX.	100.00	100.00	35.14 64.86	100.00	1	¥ X	82.68 17.32	100.00	25.27 5 7.41	6.97 10.35	100.00
	TONS/D	933.38	933.38	327.95 605.43	933.38	: :	TONS/0	49.76	60.19	15.21	4.20 6.23	60.19
S CE	LV.	100.00	100.00	39.84 60.15	100.001	; 3	Š	82.49	100.00	54.03	9.72	63.75
ERIAL BALA	68 170	8,929	8676.8	3457.2 5219.5	8676.7	RIAL BALAN	BB CO	524.6	615.9	343.6	61.8	9.209.
DETAILED MATERIAL BALANCE DEBUTANIZER COMPOSITE YIELDS	SP.GR.	.6144		.5418		DETAILED MATERIAL BALANCE DEPROPANIZER COMPOSITE YIELDS	SP.GR.	.5418		4475.	.5756	
	CHARGE	SPLIT OV LIQ=JPB-DH	CHARGE	PRODUCTS DEBUT OVER'D=JPB-DH LT NAPHTHA JP-8 DH	PRODUCTS	!	CHARGE	DEBUT OVER10=JPB-DH DEBUT OVFRHEAD-98R	CHARGE	PRODUCTS DEPROP OV.D JP-8 DH MIXED C4.5.JP-8 DH DEPROP OV.D-988 DIT	MIXED C4.8-98R PLT	PRODUCTS

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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

	P.D	03 77 35	3.2 0.1		25.7 25.8 35.8 31.8		The state of the s	۰	99			
	MSCFD	25083.83 4217.77 601.35	103.91 45.32 1977.01 79.68	i ,	414.18 24848.09 4120.27 592.53 284.53 1919.81	79.68	- 1 -	MSCFD	2186.55			
	7	53.04 36.30 3.37	1.98 17.4 88.	100.00	2.43 35.73 3.25 1.86 6.37	.55		H 73	100.001	100.00	89.38 10.62	100.001
, ,	10NS/0	406.51 278.19 25.80	15.21 41 36.08 4,20	766.39	18.61 396.47 273.82 26.94 14.43 33.52	766.39		TONS/D	96.18	98.19	87.75	96.18
	ĽĶ	100.00	:	100.00	,		8	Š				
RIAL BALAN TER _ LOS	88170	2932.6	; ;	2932.6	! !	i i	ITAL BALAN DS	28				!
DETAILED MATERIAL BALANCE FUEL GAS TREATER _ COMPOSITE YIELDS	SP.GR.	. 5418	1	F		; ; ;	DETAILED MATERIAL BALANCE SULFUR PLANT COMPOSITE YIELDS	% .			;	
	CHARGE	LP FLASH GAS: JPB-DH DEBUT OVER: D: JPB-DH SPLITTER GAS: JPB-DH DFDDAD OV: D: D: D: D:	NAP TO STAN ON STAN ON STAN ON STAN ON STAN OVER O DEPROP OV 10 - 98R PLT	CHARGE	PRODUCTS HYDROGEN SULFIDE TRI LP FLASH=JP8-DM TRI DEB OV'0=JP8-DM TRI DEP OV'0=JP8 DM TRI NAP HIRT GAS TRI DEPRO OV-948 PIT	PRODUCTS	·	CHARGE	HYDROGEN SULFIDE	CAAKGE CAAKGE BOTTA FT 6	55	PRODUCTS

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0

USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

			1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1											the transport of the control of the			and the second control of the second control							•							to the same of the
		MSCFD	24A4A 00	4120.27	F. 9.2	286.53	70 6A	74.00	45.36	1919.81	3742.77	129.46				222935.00						HSCFD		75. 28				45.32	1		
		1 4	00	20.63	; -	3 2	9 =	100	50.	2.50	2.17	. 39	41.64	100.00		53.42	46.58	100.00	1			MTZ.	8	2 2	3.79	100.00		. 12	3.79	%·30	100.00
ı		TONS/D	196.47	273.82	70.42	16.61	4.20); F	3	33.52	29.09	5.18	557.99	1150.05		715.82	624.23	1340.05	:			TONS/D	345, 31	36	13.61	359.15		.41	13.61	345.13	359.15
3		Ľ					:						100.00	- 100-00					NCE:			7/A7	97.09		2.91	100.00			2.91	97.09	100.00
HATERIAL BALANCE	YIELDS	88170											4010.5	- 4910.5					MATERIAL BALANCE	YDROTREATER	YIELDS	88170	SEAR 2		11.1	2665.9		;	77.7	2588.3	2666.0
DETAILED MATER HYDROGEN PLANT	COMPOSITE YI	SP. GR.											.6625						DETATLED MAT	•	COMPOSITE YI	SP. GR.	74.20		7.0000				1.0000	.761	
			LHANGE TOT I D ELACHE BALDH	TRI DEB OVIDE SPREAM	TOT SOLT TOTAL	DED OV. D. 108 DH	TRE DESCO ON-988 BIT	THE DEPTH OF THE	מאס ואוד קשע יאיי	TRT F.O. STAB OV'D	PLAT NET SEP GAS-988	PLAT DEBUT GAS-98R	LT NAPHTHA JP-8 DM	CHARGE	PRODUCTS	HYDROGEN (97%)	CO2HYD. PLT,	PRODUCTS					CHARGE SPITTED BOTTONS	CACO NACODAN	COLD TREATED HATER	CHARGE	PRODUCTS	NAP HYDROTREAT GAS	SOUR HATER-NAP HT	TRI HC HVY NAPHTHA	PRODUCTS

15:11 NOV 20, 61 PAGE 16			HSCFD			3742.77 129.46 204.42			MSCFD						HSCFO	2186.55		2186.55	
				_	_		_								:		•		•
			MTX	100.00	100.00	6.43 1.50 3.02 87.05	100.00		XTX	20.19 59.78 19.59	100.00	90.41 9.41 18	100.00		MTX	100.00	100.00	100.00	100.00
			10NS/0	345.13	345.13	29.09 5.18 10.42 300.44	345.13		10NS/D	623.02 1844.54 604.47 13.61	\$065.64	2789.58 290.50 5.55	3085.64		TONS/D	96.18	98.18	98.18	96.18
P51.0		BALANCE	Š	100.00	100.00	4.30 81.26	85.56	ACE	Š	20.19 59.78 19.59	100.00	90.41	90.41	BALANCE	LVX				
MIZATION REPORTS,			88.79	2568.3	2586.3	111.3	2214.5	AILED HATERIAL BALANCE R HATER TREATING POSITE YIELDS	8879	3558.4 10535.1 3452.4	17623.6	15932.7	15932.7	TERIAL BAL ERATION IELDS	88.70				
ID OPTIMIZATION	JF-2/DFH DIESEL	DETAILED MATERIAL UOP PLATFORMING COMPOSITE YIELDS	SP.GB.	. 7616		.5347		DETAILED MATERIAL B. SOUR MATER TREATING COMPOSITE YIELDS	SP. GR.	1.0000 1.0000 1.0000		1.0000		DETAILED MATERIAL (AMINE REGENERATION COMPOSITE YIELDS	SP. GR.				
UNIVERSAL OIL PRODUCTS CO. STANDARD OPTI	USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL		,	TRT HC HVY NAPHTHA	CHARGE	PRODUCTS PLAT NET SEP GAS-96R PLAT DEBUT GAS-96R DEBUT OVERHEAD-96R 96 RON PLATFORMATE	PRODUCTS	_	97-	SOUR WATER - L.P.HT SOUR WATER - H.P.HT SOUR WATER (HC) SOUR WATER-NAP HT	CHARGE	PRODUCTS TREATED SOUR WATER APPONIA LOSS	PRODUCTS			CHANGE HYDROGEN SULFIDE	CHARGE	PRODUCTS HYDROGEN SULFIDE	PRODUCTS

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14441.92 100.00 14441.92 100.00

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CHARGE OCCIDENTAL SHALE OIL

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STANDARD OPTIM	FUELS
UNIVERSAL OIL PRODUCTS CO.	USAF SHALE OIL TO FUELS
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CASE 4 - JP-8 JET FUEL + DF-2/DFH DIESEL

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MSCFD				;		MSCFD			1977.01	Managhan in a crimination of the contract of t		MSCFD
Z Z	100.00	100.00	5.03	100.00		MTX	100.00	100.00	1.82	300.00	!	77.
10NS/D	290.50	290.50	14.62	290.50	;	TONS/D	1985.51	1965.51	36.08	1985.51		TONS/D
Ĕ					3	7	100.00	100.00	95.19	95.19	; ;	ĘĶ
88170			83.5	83.5	RIAL BALAN MILIZER LOS	88C0	13537.4	13537.4	12886.8	12886.8	RIAL BALAN LITIES	88170
SP. GR.			1.0000	1	DETAILED MATE FUEL DIL STAE COMPOSITE YIE	SP. GR.	7780.		0498.		COMBINED FACT	SP.GR.
CHARGE	AITIONIA	CHARGE	PRODUCTS TREATED SOUR WATER ANHYDRUS APPONIA	PRODUCTS			SEPARATOR LIQUID(HP)	CHARGE	PRODUCTS F.O. STAB OVER'D F.O. STAB BOTTOMS	PRODUCTS		CHARGE
	SP.GR. BBL/D LVX TONS/D MTX	SP.GR. BBL/D LVX TONS/D HTX HSCFD VPTONIA 290.50 100.00	SP.GR. BBL/D LV% TONS/D HT% HSCFD UPTONIA 290.50 100.00 CHARGE 290.50 100.00	SP.GR. BBL/D LV.: TONS/D HT.: HSCFD 290.50 100.00 290.50 100.00 290.50 100.00 1.0000 83.5 14.62 5.03 275.86 94.97	SP.GR. BBL/D LV% TONS/D WT% HSCFD 290.50 100.00 290.50 100.00 290.50 100.00 275.88 94.97 275.88 94.97	SP.GR. BBL/D LV.: TONS/D WTY. HSCFD 290.50 100.00 290.50 100.00 290.50 100.00 33.5 275.88 94.97 83.5 290.50 100.00 DETAILED MATERIAL BALANCE FUEL DIL STABILIZER COMPOSITE YIELDS	SP. GR. BBL/D LV% TONS/D WTY HSCFD CHARGE CHARGE TS TREATED SOLR WATER 1.0000 83.5 TREATED SOLR WATER 1.0000 83.5 TREATED SOLR WATER 1.0000 83.5 275.88 94.97 PRODUCTS DETAILED HATERIAL BALANCE FUEL DIL STABILIZER CCHPOSITE YIELDS SP. GR. BBL/D LV% TONS/D MT% MSCFD	CHARGE ATTONIA CHARGE CHARGE PRODUCTS TREATED SOLR HATER ANTONIOCTS TREATED SOLR HATER ANTORIUS ATTONIA PRODUCTS CHARGE	SP.GR. BBL/D LVX TONS/D HTX HSCFD LONG/D HTX HS	CHARGE APTONIA CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE CHARGE SEPARATOR LIGUIDIHP) CHARGE F.O. STAB BOYER D F.O. STAB BOYER D F.O. STAB BOYER D F.O. STAB BOYER D F.O. STAB BOTGHS CHARGE APTONIC STAB BOYER D F.O. STAB BOTGHS CHARGE APTONIC SPORT S CHARGE APTONIC SPORT S CHARGE APTONIC SPORT S CHARGE APTONIC SPORT S CHARGE APTONIC SPORT S F.O. STAB BOYER D F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTGHS F.O. STAB BOTTGHS GR. BBL/D LVZ TOKS/D HTZ HSCFD	CHARGE PRODUCTS THEATED SOLE HATER THEATER	

LATION REPORTS, PS1.0] ESEL	DETAILED HATERIAL BALANCE DUTHY LINIT		SP.GR. BBL/D LV% TONS/D HT%	.5746 405.4 100.00 40.78 100.00	405.4 100.00 40.78 100.00	.5743 405.6 100.05 40.78 100.00	
UNIVERSAL DIL PRODUCTS CO. STANDARD OPTIHIZATION REPORTS, PS1.0	USAF SHALE DIL TO FUELS PHASE IV	CASE 4 - JP-8 JET FUEL + DF-2/DFH DIESEL	DETAILED HA	COHOS		MIXED BUTANES	CHARGE	PRODUCTS PIXED BUTANES	BOTHITE

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0 USAF SHALE OIL TO FUELS PHASE IV

DIESEL	
0F-2/DFH	
r FUEL 1	
JP-8 JET	
CASE 4	

		DETAILED USAGE OF AN UNPOOLED UTILITY ELECTRIC POHER UNIT IS KWH COSTS ARE ALLOCATED TO PROCESS UNITS	N UNPOOLED UTILITY IT IS KNH TO PROCESS UNITS	Y PRICE IS	.04500/UNIT) !		1
PROCESS UNIT	HOOE	CONTROL STREAM	8879	TONS/D	UNIT/BBL	UNITATON	UNITS/D	9,
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	90000.0	14441.92	0801	144	97.00 AE	44 614
L.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	90000	14441.92	1.9070	11.886	171627 72	7721 26
H.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.0	14390,21	1.2677	A. 160	117422 70	K286.02
HYDROCRACKING	COMBINED MODES	COMPOSITE USAGE	84889.1	12450.46	6.1047	41.622	518213.05	21110.50
NAPHINA SPLITTER	COMBINED MODES	COMPOSITE USAGE	11848.5	1338.05	3912	3.464	4634.74	208.56
DEBUTANIZER	COMBINED MODES		8676.8	933.38	.1399	1,301	1213.96	54.65
OF PROPANIZER	COMBINED MODES		635.9	60.19	1200	1.268	76. 12	7
FUEL GAS TREATER	COMBINED MODES			766.39			•	
SULFUR PLANT	COMBINED MODES		-	98.18		21,974	2157.44	97 CB
HIDROGEN PLANT	COMBINED MODES			715.82		211.850	151646 10	A824 07
NAPHTHA HYDROTREATER	COMBINED MODES	COMPOSITE USAGE	2588.2	345.31	. 7065	5.2%	1828.60	B2.29
OUP PLATFORMING	COMBINED MODES		2500.3	345.13	2,3309	17.680	6032.93	271.68
SOUR MATER TREATING	COMBINED MODES	COMPOSITE USAGE	17623.6	3085.64	. 3787	2.163	60.429	300.33
ATIME REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.18		40.773	4003.15	180.16
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.88		101	0.45	40.10
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	13537.4	1985.51	1638	1.116	2216 83	2
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000	14441.92				
DUTY UNIT	COMBINED MODES	-	4.05.4	40. 7A				

UNIVERSAL OIL PRODUCTS CO.		STANDARD OPTIMIZATION REPORTS, PS1.0	P51.0			15:11	15:11 NOV 20, 61	PAGE 20	
USAF SHALE PHASE IV	USAF SHALE OIL TO FUELS PHASE IV								
CASE 4 - JF	CASE 4 - JP-8 JET FUEL + DF-2/	-2/DFH DIESEL							
		DETAILED USAGE OF A POOLED UTILITY 600# STEAM UNIT IS MLBS COSTS ADE ALLOCATED TO LITTLE AND	COLED UTILITY 15 HLBS					4	
PROCESS UNIT	HODE	CONTROL STREAM		TONS/0	UNITZBBL	UNITATON	UNITS/D	\$	
			• • •					;	
FEED PREPARATION		COMPOSITE USAGE	90000.0	14441.92			:	•	
L.P. MYDROTREATING	_	_	90000.0	14441.92	.0193	. 120	1733.03		
HADDONDACKATING	_	_	92629.0	14390.21	.0158	. 102	1462.05		
MICHACKING MICHAEL	_	_	64889.1	12450.46		******		-	
DEPLITA SPLITER	_	COMPOSITE USAGE	11848.5	1338.05					
DEBOT ANI ZER	_	COMPOSITE USAGE	8,926.8	933.38					
DE PROPANI ZER		COMPOSITE USAGE	635.9	60.19					
FUEL GAS TREATER	_	COMPOSITE USAGE		766.39			!		
SULFUR PLANT	_	COMPOSITE USAGE		96.18		. 220	21.57		
HIDROGEN PLANT	COMBINED MODES	COMPOSITE USAGE		715.82		-6.696	-4791.42		
NAPHTHA HYDROTREATER		COMPOSITE USAGE	2588.2	345.31			!		
UOP PLATFORMING		COMPOSITE USAGE	2588.3	345.13	-,0603	452	-155.97		
SOUR MATER TREATING	COMBINED MODES	COMPOSITE USAGE	17623.6	3065.64					
AMINE REGENERATION	-	COMPOSITE USAGE	•	99.18					
APPONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.88					
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	13537.4	1985.51					
COMBINED FACILITIES	COMBINED MODES	_	90000	14441.92			: :		
burn unit	COMBINED MODES	COMPOSITE USAGE	405.4	40.78					
TOTALS			; ; ;		į		-1730,74		
									i
ANOUNT PRODUCED FROM REFINERY FUELS	REFINERY FUELS						754.67	514.55	
ANDLINT CONVERTED TO 150# STEAM	50# STEAM	1	•				2485.42		

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

	;	DETAILED USAGE OF COOLING WATER COSTS ARE ALLOCAT	DETAILED USAGE OF AN UNPOOLED UTILITY COOLING HATER UNIT IS MGAL COSTS ARE ALLOCATED TO PROCESS UNITS	PRICE IS	03000/WIT.		i i	:
PROCESS UNIT	700E	CONTROL STREAM	BB/70	10NS/D	UNITABBL	UNITZION	QVITS/D	9/9
FEED PREPARATION	COMBINED MODE	S COMPOSITE USAGE	0.0000.0	14441.92		***************************************		:
L.P. HYDROTREATING	COMBINED MODE	COMPOSITE	90000.0	14441.92	.0016	.010	145.86	2.
H.P. HYDROTREATING	COMBINED MODE	COMPOSITE	92629.0	14390.21				
HYDROCRACKING	COMBINED MODE	S COMPOSITE USAGE.	64868.1	12450.46	. 0263	179	2228.63	8.8
NAPHTHA SPLITTER	COMBINED MODES	COMPOSITE	11848.5	1338.05	.0299	.264	353.91	10.62
DEBUTANI ZER	COMBINED MODE	COMPOSITE	8676.8	933.38	.0116	. 107	100.34	3.01
DEPROPAVIZER	COMBINED MODE	COMPOSITE	635.9	60.19	.0857	. 905	54.50	1.63
FUEL GAS TREATER	COMBINED MODE	΄,		766.39			; ; ;	:
SULFUR PLANT	COMBINED MODE	COMPOSITE		98.18				
HYDROGEN PLANT	COMBINED MODE	COMPOSITE		715.82		.535	362.75	11.46
NAPHTHA HYDROTREATER	COMBINED MODE	S COMPOSITE USAGE	2586.2	345.31	.0118	.069	30.66	. 92
UOP PLATFORMING	COMBINED MODE	COMPOSITE	2588.3	345.13	0800	909	207.08	6.21
SOUR MATER TREATING	COMBINED MODE	COMPOSITE	17623.6	3085.64	. 1336	. 763	2354.64	70.64
AMINE REGENERATION	COMBINED MODE	COMPOSITE		98.18		32.389	3180.00	95.40
AMIONIA PLANT	COMBINED MODE	COMPOSITE		275.88		10.%1	3023.94	90.72
FUEL OIL STABILIZER	COMBINED MODE	COMPOSITE	13537.9	1985.51	.0200	137	271.62	9.14
COMBINED FACILITIES	COMBINED HODE	S COMPOSITE USAGE	90000.0	14441.92				
DUTY UNIT	COMBINED HODE	S COMPOSITE USAGE	402.4	£0.3				

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REPORTS,
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USAF SHALE DIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

FEED FREPARATION L.P. HYDROTREATING H.P. HYDROTREATING HYDROCRACKING NAPHTHA SPLITTER DEBUTANIZER DEPROPANIZER FUEL GAS TREATER SULFUR PLANT NAPHTHA HYDROTREATER UOP PLATFORMING SOUR MATER TREATING AMPINA BI ANT	COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES COPBINED HODES	COMPOSITE USAGE COMPOSITE USAG	REFINERY FUELS UNIT IS HEBTU COSTS ARE ALLOCATED TO UTILITY POOL COMPOSITE USAGE COMPOSITE USA	14441.92 14441.92 14590.21 12550.46 1318.05 933.38 60.19 766.39 98.18 345.31 365.64	. 0218 . 04.62 . 1721 . 0640 . 3096	 UNITS/D 1961.21 4665.72 16609.37 756.67 43.96 43.92	
FUEL OIL STABILIZER COMBINED FACILITIES DUMNY UNIT		COMPOSITE USAGE COMPOSITE USAGE COMPOSITE USAGE	13537.4 90000.0 405.4	275.88 1985.51 14441.92 40.78	9850.	 . 658,20	

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496.07

68425.25

67739.19

ANDUNT CONVERTED TO 600# STEAN

AHOUNT PRODUCED FROM F.O. STAB BOTTOMS

TOTALS

STANDARD OPTIMIZATION REPORTS, P51.0 UNIVERSAL OIL PRODUCTS CO.

		DETAILED USAGE OF A POOLED UTILITY 150# STEAM UNIT 1S HLBS COSTS ARE ALLOCATED TO UTILITY POOL	ED UTILITY MLBS TILITY POOL	,	; ;	1	:	
PROCESS UNIT	700E	CONTROL STREAM	BB/70	TONS/D	UNIT/88L	UNITATON	DNITS/D	0/*
FEED PREPARATION	COMBINED HODES	COMPOSITE USAGE	90000.0	14441.92	!			
L.P. HYDROTREATING	COMBINED MODES	_	90000	14441.92				
H.P. HYDROTREATING		COMPOSITE USAGE	92629.0	14390.21				
HYDROCRACKING		_	84688.1	12450.46	0185	#.126	#.126 -1573.74	
NAPHTHA SPLITTER		_	11648.5	1338.05				
DEBUTANI ZER		_	8676.8	933.38				
DEPROPANI ZER	COMBINED MODES	_	635.9	60.19				
FUEL GAS TREATER	COMBINED MODES		:	766,39	1	1		
SULFUR PLANT	COMBINED MODES	COMPOSITE USAGE		98.18	<u>.</u>	-7.593	-745.50	
HYDROGEN PLANT	COMBINED MODES			715.82				
NAPHTHA HYDROTREATER		_	2588.2	345.31				
UOP PLATFORMING		COMPOSITE USAGE	2588.3	345.13	7000	200	1.79	
SOUR MATER TREATING		COMPOSITE USAGE	17623.6	3085.64	. 2633	1.504	4639.86	
AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		96.18		1.660	163.00	
APPIONIA PLANT	COMBINED MODES	COMPOSITE USAGE		275.88				
		_	13537.4	1995.51				
COMBINED FACILITIES	_	_	90000	14441.92				
LIN JUNO 4	COMBINED HODES	COMPOSITE USAGE	405.4	40.78				
TOTALS	-						2485.42	
ATOUNT PRODUCED FROM 600# STEAM	600# STEAM						2485.42	

Appendix B.8 (Cont.)

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STANDARD OPTIMIZATION REPORTS, PS1.0 UNIVERSAL OIL PRODUCTS CO.

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		OF-2/OFM DIESEL
s		٠
TO FUELS		FUEL
2		JET
_		
OIL		9-9
		5
SHALE	>	•
Ď		4
USAF	PHASE	CASE 4

	UNITS/D \$/D	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			-040 A1		100 21	17 55		17 676	7				01 20				
	UNITATON				070 -		869	200	•	778 0-	3				0 127				
	UNIT/BBL				7110		0460	0276						:				:	
	10NS/0	14441.92	14441.92	14390.21	12450.46	1338.05	953.38	60.19	766.39	98.18	715.82	345.31	345.13	3085.64	98.18	275.88		14441.92	,
SAGE OF A POOLED UTILITY LINIT IS MLBS ALLOCATED TO UTILITY POOL	88 Vo	90000.0	90000	92629.0	84888.1	11848.5	8676.8	632.9				2588.2	2588.3	17623.6			13537.4	90000	A 50.0
DETAILED U 50# STEAM- COSTS ARE	CONTROL STREAM	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	_	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	_	_	-			COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSITE USAGE	COMPOSTAT I MAGE
	MODE						COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED	COMBINED	COMBINED, MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES	COMBINED MODES
	PROCESS UNIT	FEED PREPARATION	L. P. HTOROTREATING	H. P. HYDROTREATING	HOROCRACKING	NAPHTHA SPLITTER	DEBUTANI ZER	DEPROPANI ZER	FUEL GAS TREATER	SULFUR PLANT	HYDROGEN PLANT	NAPHTHA HYDROTREATER	UOP PLATFORMING	SOUR MATER TREATING	AMINE REGENERATION	ATTONIA PLANT	FUEL OIL STABILIZER	CUTBINED FACILITIES	

TOTALS

- 2973.74

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIHIZATION REPORTS, PS1.0 USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-6 JET FUEL + 0F-2/DFH DIESEL

		DETAILED USAGE OF A POOLED UTILITY BOILER WATER UNIT IS MLBS FORTS ARE ALL MONTED TO LITTLE TO DO	LED UTILITY S MLBS					; i
Ī	HOOE	CONTROL STREAM		1015/0	INTTABA	NOTATIVE	2	•
Ū	COMBINED MODES	no say attached						1
ב	COMBINED MODES		90000	26 Tabat				
1			90000.0	14441.92				
3	CTBINED MODES	COMPOSITE USAGE	92629.0	14390.21				
ັ	OFBINED MODES	COMPOSITE USAGE	84886.1	12450.46	0610	266	12 4127	
ັ	OFBINED MODES	COMPOSITE USAGE	11848.5	1338.05				
ŭ	COMBINED MODES	COMPOSITE USAGE	8676.8	933.38				
ၓ	OMBINED MODES	COMPOSITE USAGE	632.9	60.19				
ŭ	OMBINED MODES	COMPOSITE USAGE		766.39				
ັບ	COMBINED MODES	COMPOSITE USAGE		98.18	!	18 066	1840 15	1 1
ຽ	OMBINED MODES	COMPOSITE USAGE		715.82		10, 01	11860 44	
ၓ	OMBINED MODES	COMPOSITE USAGE	2588.2	145.31				
ၓ	OMBINED MODES	COMPOSITE USAGE	2568.3	345.13	0100	704	26.9 64	
ដ	OPPINED MODES	COMPOSITE USAGE	17623.6	3085.64	1		The Tales	
Ծ		COMPOSITE USAGE		8				
ដ	COMBINED MODES	COMPOSITE USAGE		275.88				
ដ	COMBINED MODES	COMPOSITE USAGE	13537.4	1985, 51				
Ծ	COMBINED MODES	COMPOSITE USAGE	0.00006	14441 02	:			
8	COMBINED MODES	COMPOSITE USAGE	402.4	40.78				
				•	:		19297. 66	
								!
							9824.12	4912.06

AMOUNT PRODUCED FROM CONDENSATE

9473.74

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USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

		DETAILED USAGE OF A POOLED UTILITY CONDENSATE UNIT IS MUSS	ED UTILITY				
		ALLOCAT	JUILITY POOL				
PROCESS UNIT	MODE	CONTROL STREAM	88L/0	TONS/D	UNIT/BBL	UNIT/TON	UNITS/D
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	90000	14441.92		;	
L.P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	90000	14441.92	0193	-, 120	-1733.03
H. P. HYDROTREATING	COMBINED MODES	COMPOSITE USAGE	92629.0	14390.21	0158	-, 102	-1462.05
HYDROCHACKING	_	COMPOSITE USAGE	84888.1	12450.46			
NAPHINA SPLITTER	_	COMPOSITE USAGE	11648.5	1338.05			
DEBUTANIZER	U	COMPOSITE USAGE	8676.8	933.38	0460	428	-399.21
DE PROPANI ZER	U	COMPOSITE USAGE	635.9	60.19	0276	-, 292	-17.55
FUEL GAS TREATER	U	COMPOSITE USAGE		766.39) i	•	
SULFUR PLANT	J	COMPOSITE USAGE		96.18		684	-67.12
HYDROGEN PLANT	U	COMPOSITE USAGE		715.82		•	
NAPHTHA HYDROTREATER	J	COMPOSITE USAGE	2586.2	345.31			
UOP PLATFORMING	COMBINED	COMPOSITE USAGE	2588.3	345.13	-,0294	221	-76.21
SOUR MATER TREATING	COMBINED MODES	COMPOSITE USAGE	17623.6	3085.64	2633	-1.504	-64 10 BA
AMINE REGENERATION	COMBINED MODES	COMPOSITE USAGE		98.18		700 01	-1076 72
APPONIA PLANT	COMBINED MODES			275.88			
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE USAGE	13537.4	1985.51			
COMBINED FACILITIES	COMBINED MODES	COMPOSITE USAGE	90000	14441.92	:		1
DUTY UNIT	COMBINED MODES	COMPOSITE USAGE	405.4	40.78			

AHOUNT CONVERTED TO BOILER WATER

UNIVERSAL OIL PRODUCTS CO. STANDARD OPTIMIZATION REPORTS, P51.0
USAF SHALE OIL TO FUELS
PHASE IV
CASE 4 - JF-8 JET FUEL + DF-2/DFM DIESEL

		DETAILED USAGE OF AN UNPOOLED UTILITY CAT. & CHEMICALS UNIT IS \$	NPOOLED UTILITY		PRICE IS 1.00000/UNIT			;
		COSTS ARE ALLOCATED TO PROCESS UNITS	PROCESS UNITS					
PROCESS UNIT	700E	CONTROL STREAM	B8L/D	TONS/D	UNIT/BBL	UNITATON	UNITS/0	9,4
FEED PREPARATION	COMBINED MODES	COMPOSITE USAGE	0.00006	14441.92		•	:	
L.P. HIDROTREATING	COMBINED MODES	COMPOSITE	90000.0	14441.92	.1145	.714	10308.64	10309.64
M.P. HYDROTREATING	COMBINED MODES	COMPOSITE	92629.0	14390.21	.0417	. 268	3658.02	3858.02
HYDROCRACKING	COMBINED MODES	COMPOSITE USAGE	84888.1	12450.46	.057A	394	4910.66	4910.46
NAPHTHA SPLITTER	COMBINED MODES	COMPOSITE	11848.5	1338.05				
DEBUT AVI ZER	COMBINED MODES	COMPOSITE USAGE	8676.8	933.38				
DEPROPAUI ZER	COMBINED MODES	COMPOSITE USAGE	635.9	60.19				
FUEL GAS TPEATER	COMBINED MODES	COMPOSITE USAGE		766.39	:		1	i
SULFUR PLANT	CUMBINED MODES	COMPOSITE USAGE		98.18				
MICROGEN PLANT		COMPOSITE USAGE		715.82		5.191	3715.56	3715.56
NAPHTHA HYDROTREATER		Ü	2588.2	345.31	.0028	.021	7.36	7.36
UOP PLATFORMING	COMBINED MODES	_	2589.3	345, 13	5640	.326	112,55	112.55
SOUR HATER TREATING	COMBINED MODES	COMPOSITE	17623.6	3085.64				
AMINE REGENERATION	COMBINED MODES	Ĭ		96.18		. 283	27.75	27.75
APPONIA PLANT	COMBINED MODES	Ö		275.89				
FUEL OIL STABILIZER	COMBINED MODES	COMPOSITE	13537.4	1935.51			i	į
COMBINED FACILITIES	COMBINED MODES	COMPOSITE	900006	14441.92			 - -	
חשיו האז האז	COMBINED MODES	COMPOSITE	4 504	40. 7A				

TOTALS

USAF SMALE OIL TO FUELS PMASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL

PROCESS UNIT UTILITY CONSUMPTION SUFFARY

מחרכים פיני	ELECTRIC POWER KWH	600# STE AM MLBS	COOLING HATER HGAL	REFINERY FUELS MPBTU	150# STE AM MLBS	50# STEA M MLBS	BOILER H ATER MLBS	CONDENSA TE MLBS
FEED PREPARATION	9720.9							
L. P. HYDROTREATING	171627.7	1733.0	145.9	1961.2				-1733.0
HYDROTREATING	117422.7	1462.0		4466.7				-1462.0
NOROCRACKING	518213.0	!	2228.6	14609.4	-1573.7	-989.B	1316.3	i :
VAPHTHA SPLITTER	4634.7		353.9	7.58.7				
DEBUTANI ZER	1214.0		100.3			399.2		-399.2
DE PROPANI ZER	76.3		54.5			17.6		-17.6
SULFUR PLANT	2157.4	21.6			-745.5	4.896	1860.1	-67.1
HOROGEN PLANT	151646.1	4.1614-	382.8	44439.7			13880.5	
WAPHTHA HYDROTREATER	1828.6		30.7	43.9				
UOP PLATFORMING	6032.9	-156.0	207.1	801.4	1.8		242.9	-76.2
MATER TREATING	6674.1		2354.6		4639.9			-4639.9
WINE REGENERATION	4003.2		3180.0		163.0	915.7		-1078.7
AMMONIA PLANT	936.0		3023.9			•		
OIL STABILIZER	2216.8		271.4	658.2				
TOTAL CONSUMPTION	998404.5	-1730.7	12333.7	67739.2	2485.4	-625.8	19297.9	-9473.7

STANDARD OPTIMIZATION REPORTS, P51.0 USAF SHALE OIL TO FUELS PHASE IV CASE 4 - JP-8 JET FUEL + DF-2/DFM DIESEL UNIVERSAL OIL PRODUCTS CO.

PROCESS UNIT UTILITY CONSUMPTION SUFFARY

HEMICALS	10308.6 3858.0	4910.5	3715.6	27.7
	FEED PREPARATION L.P. HYDROTREATING H.P. HYDROTREATING	HYDROCKACKING NAPHTHA SPLITTER DEBUTANIZER DEFROPANIZER SHEED SHAFT	HYDEGER PLANT NAMINA HYDROTREATER UDP PLATFORHING	SOUR HATER TREATING AMINE REGENERATION AMONIA PLANT FUEL OIL STABILIZER

TOTAL CONSUMPTION

DIFIC